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(71) Applicant (for all designated States except US): BIOCHEM VACCINS INC. [CA/CA]: 2323 boulevard du Parc Tech-

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(71) Applicant (for all designated States except US): BIOCHEM VACCINS INC. [CA/CA]; 2323 boulevard du Parc Technologique, Sainte-Foy, Québec G1P 4R8 (CA).

(72) Inventors; and
(75) Inventors/Applicants (for US only): BRODEUR, Bernard, R. [CA/CA]; 2401 rue Maritain, Sillery, Québec G1T 1N6 (CA). RIOUX, Clément [CA/CA]; 1012 Jean-Charles Cantin, Ville de Cap Rouge, Québec G1Y 2X1 (CA). BOYER, Martine [CA/CA]; Apt. 204, 25 des Mouettes, Beauport, Québec G1E 7G1 (CA). CHARLEBOIS, Isabelle [CA/CA]; 410 Mirabel, St-Nicolas, Québec G7A 2L5 (CA). HAMEL, Josée [CA/CA]; 2401 rue Maritain, Sillery, Québec G1T 1N6 (CA). MARTIN, Denis [CA/CA]; 4728-G rue Gaboury, St-Augustin-de-Desmaures, Québec G3A 1E9 (CA).

(74) Agents: CÔTE, France et al.; Swabey Ogilvy Renault, Suite 1600, 1981 McGill College Avenue, Montréal, Québec H3A 2Y3 (CA).

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(54) Title: GROUP B STREPTOCOCCUS ANTIGENS

(57) Abstract

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Group B streptococcus (GBS) proteins and polynucleotides encoding them are disclosed. Said proteins are antigenic and therefore useful vaccine components for the prophylaxis or therapy of streptococcus infection in animals. Also disclosed are recombinant methods of producing the protein antigens as well as diagnostic assays for detecting streptococcus bacterial infection.

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GROUP B STREPTOCOCCUS ANTIGENS

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FIELD OF THE INVENTION

The present invention is related to antigens, more particularly protein antigens of group B streptococcus (GBS) bacterial pathogen which are useful as vaccine components for therapy and/or prophylaxis.

BACKGROUND OF THE INVENTION

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Streptococcus are gram (+) bacteria that are differentiated by group specific carbohydrate antigens A through O found on their cell surface. Streptococcus groups are further distinguished by type-specific capsular polysaccharide antigens. Several serotypes have been identified for the Group B streptococcus (GBS): Ia, Ib, II, III, IV, V, VI, VII and VIII. GBS also contains antigenic proteins known as "C-proteins" (alpha, beta, gamma and delta), some of which have been cloned.

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Although GBS is a common component of the normal human vaginal and colonic flora this pathogen has long been recognized as a major cause of neonatal sepsis and meningitis, late-onset meningitis in infants, postpartum endometritis as well as mastitis in dairy herds. Expectant mothers exposed to GBS are at risk of postpartum infection and may transfer the infection to their baby as the child passes through the birth canal. Although the organism is sensitive to antibiotics, the high attack rate and rapid onset of sepsis in neonates and meningitis in infants results in high morbidity and mortality.

To find a vaccine that will protect individuals from GBS infection, researches have turned to the type-specific antigens. Unfortunately these polysaccharides have proven to be poorly immunogenic in humans and are restricted to the particular serotype from which the polysaccharide originates. Further, capsular polysaccharide elicit a T cell independent response i.e. no IgG production.

Consequently capsular polysaccharide antigens are unsuitable as a vaccine component for protection against GBS infection.

Others have focused on the C-protein beta antigen which demonstrated immunogenic properties in mice and rabbit models. This protein was found to be unsuitable as a human vaccine because of its undesirable property of interacting with high affinity and in a non-immunogenic manner with the Fc region of human IgA. The C-protein alpha antigen is rare in type III serotypes of GBS which is the serotype responsible for most GBS mediated conditions and is therefore of little use as a vaccine component.

Therefore there remains an unmet need for GBS antigens that may be used as vaccine components for the prophylaxis and/or therapy of GBS infection.

SUMMARY OF THE INVENTION

According to one aspect, the present invention provides an isolated polynucleotide encoding a polypeptide having at least 70% identity to a second polypeptide comprising a sequence selected from the group consisting of:

SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,

SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10,

SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15,

SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19,

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SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.
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In other aspects, there is provided vectors comprising polynucleotides of the invention operably linked to an expression control region, as well as host cells transfected with said vectors and methods of producing polypeptides comprising culturing said host cells under conditions suitable for expression.

In yet another aspect, there is provided novel polypeptides encoded by polynucleotides of the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

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Figure 1a is the DNA sequence of clone 1 (SEQ ID NO :1) with corresponding amino acid sequences for open reading frames; figure 1b is the amino acid sequence SEQ ID NO: 2; figure 1c is the amino acid sequence SEQ ID NO: 3; figure 1d is the amino acid sequence SEQ ID NO: 4; figure 1e is the amino acid sequence SEQ ID NO: 5; figure 1f is the amino acid sequence SEQ ID NO: 6;
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Figure 2a is the DNA sequence of clone 2 (SEQ ID NO :7) with corresponding amino acid sequences for open reading frames; figure 2b is the amino acid sequence SEQ ID NO: 8; figure 2c is the amino acid sequence SEQ ID NO: 9; figure 2d is the amino acid sequence SEQ ID NO:10; figure 2e is the amino acid sequence SEQ ID NO:11; figure 2f is the amino acid sequence SEQ ID NO:12;

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Figure 3a is the DNA sequence of clone 3 (SEQ ID NO :13)
    with corresponding amino acid sequences for open reading
    frames;
    figure 3b is the amino acid sequence SEQ ID NO:14;
    figure 3c is the amino acid sequence SEQ ID NO:15;
    figure 3d is the amino acid sequence SEQ ID NO:16;
    figure 3e is the amino acid sequence SEQ ID NO:17;
    figure 3f is the amino acid sequence SEQ ID NO:18;
    figure 3g is the amino acid sequence SEQ ID NO:19;
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    figure 3h is the amino acid sequence SEQ ID NO:20;
    figure 3i is the amino acid sequence SEQ ID NO:21;
    Figure 4a is the DNA sequence of clone 4 (SEQ ID NO :22)
    with corresponding amino acid sequences for open reading
    frames;
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    figure 4b is the amino acid sequence SEQ ID NO:23;
    figure 4c is the amino acid sequence SEQ ID NO:24;
    figure 4d is the amino acid sequence SEQ ID NO:25;
    figure 4e is the amino acid sequence SEQ ID NO:26;
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    Figure 5a is the DNA sequence of clone 5 (SEQ ID NO :27)
    with corresponding amino acid sequences for open reading
    frames;
    figure 5b is the amino acid sequence SEQ ID NO:28;
    figure 5c is the amino acid sequence SEQ ID NO:29;
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    figure 5d is the amino acid sequence SEQ ID NO:30;
    figure 5e is the amino acid sequence SEQ ID NO:31;
    Figure 6a is the DNA sequence of clone 6 (SEQ ID NO :32) ;
    figure 6b is the amino acid sequence SEQ ID NO:33;
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    figure 6c is the amino acid sequence SEQ ID NO:34;
    figure 6d is the amino acid sequence SEQ ID NO:35;
    figure 6e is the amino acid sequence SEQ ID NO:36;
    Figure 7a is the DNA sequence of clone 7 (SEQ ID NO :37);
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    figure 7b is the amino acid sequence SEQ ID NO:38;
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figure 7c is the amino acid sequence SEQ ID NO:39; figure 7d is the amino acid sequence SEQ ID NO:40; figure 7e is the amino acid sequence SEQ ID NO:41;
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Figure 8 is the DNA sequence of a part of clone 7 including a signal sequence (SEQ ID NO :42);

Figure 9 is the DNA sequence of a part of clone 7 without a signal sequence (SEQ ID NO :43);

10 Figure 9a is the amino acid sequence (SEQ ID NO:44);

Figure 10 represents the distribution of anti-GBS ELISA titers in sera from CD-1 mice immunized with recombinant GBS protein corresponding to the SEQ ID NO:39.

DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to novel antigenic polypeptides of group B streptococcus (GBS) characterized by the amino acid sequence selected from the group consisting of:

SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,

SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10,

SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15,

10 SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19,

SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24,

SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29,

SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34,

SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39,

15 SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.

A preferred embodiment of the invention includes SEQ ID NO:39 and SEQ ID NO:44.

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A further preferred embodiment of the invention is SEQ ID NO :39.

A further preferred embodiment of the invention is SEQ ID NO :44.

As used herein, "fragments", "derivatives" or "analogs" of the polypeptides of the invention include those polypeptides in which one or more of the amino acid residues are substituted with a conserved or non-conserved amino acid residue (preferably conserved) and which may be natural or unnatural.

The terms «fragments», «derivatives» or «analogues» of polypeptides of the present invention also include polypeptides which are modified by addition, deletion,

substitution of amino acids provided that the polypeptides retain the capacity to induce an immune response.

By the term «conserved amino acid» is meant a substitution of one or more amino acids for another in which the antigenic determinant (including its secondary structure and hydropathic nature) of a given antigen is completely or partially conserved in spite of the substitution.

- 10 For example, one or more amino acid residues within the sequence can be substituted by another amino acid of a similar polarity, which acts as a functional equivalent, resulting in a silent alteration. Substitutes for an amino acid within the sequence may be selected from other members 15 of the class to which the amino acid belongs. For example, the nonpolar (hydrophobic) amino acids include alanine, leucine, isoleucine, valine, proline, phenylalanine, tryptophan and methionine. The polar neutral amino acids include glycine, serine, threonine, cysteine, tyrosine, 20 asparagine and glutamine. The positively charged (basic) amino acids include arginine, lysine and histidine. The negatively charged (acidic) amino acids include aspartic acid and glutamic acid.
- 25 Preferably, derivatives and analogs of polypeptides of the invention will have about 70% identity with those sequences illustrated in the figures or fragments thereof. That is, 70% of the residues are the same. More preferably polypeptides will have greater than 95% homology. In another preferred embodiment, derivatives and analogs of polypeptides of the invention will have fewer than about 20 amino acid residue substitutions, modifications or deletions and more preferably less than 10. Preferred substitutions are those known in the art as conserved i.e. the substituted residues share physical or chemical properties such as hydrophobicity, size, charge or functional groups.

Furthermore, in those situations where amino acid regions are found to be polymorphic, it may be desirable to vary one or more particular amino acids to more effectively mimic the different epitopes of the different GBS strains.

Also included are polypeptides which have fused thereto other compounds which alter the polypeptides biological or pharmacological properties i.e. polyethylene glycol (PEG) to increase half-life; leader or secretory amino acid sequences for ease of purification; prepro- and pro- sequences; and (poly) saccharides.

Moreover, the polypeptides of the present invention can be modified by terminal -NH₂ acylation (eg. by acetylation, or thioglycolic acid amidation, terminal carbosy amidation, e.g. with ammonia or methylamine) to provide stability, increased hydrophobicity for linking or binding to a support or other molecule.

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Also contemplated are hetero and homo polypeptide multimers of the polypeptide fragments, analogues and derivatives. These polymeric forms include, for example, one or more polypeptides that have been cross-linked with cross-linkers such as avidin/biotin, gluteraldehyde or dimethyl-superimidate. Such polymeric forms also include polypeptides containing two or more tandem or inverted contiguous sequences, produced from multicistronic mRNAs generated by recombinant DNA technology.

Preferably, a fragment, analog or derivative of a polypeptide of the invention will comprise at least one antigenic region i.e. at least one epitope.

In order to achieve the formation of antigenic polymers

(i.e. synthetic multimers), polypeptides may be utilized having bishaloacetyl groups, nitroarylhalides, or the like,

where the reagents being specific for thio groups. Therefore, the link between two mercapto groups of the different peptides may be a single bond or may be composed of a linking group of at least two, typically at least four, and not more than 16, but usually not more than about 14 carbon atoms.

In a particular embodiment, polypeptide fragments, analogs and derivatives of the invention do not contain a methionine (Met) starting residue. Preferably, polypeptides will not incorporate a leader or secretory sequence (signal sequence). The signal portion of a polypeptide of the invention may be determined according to established molecular biological techniques. In general, the polypeptide of interest may be isolated from a GBS culture and subsequently sequenced to determine the initial residue of the mature protein and therefor the sequence of the mature polypeptide.

- According to another aspect, there is provided vaccine compositions comprising one or more GBS polypeptides of the invention in admixture with a pharmaceutically acceptable carrier diluent or adjuvant.
- Suitable adjuvants include oils i.e. Freund's complete or incomplete adjuvant; salts i.e. AlK(SO₄)₂, AlNa(SO₄)₂, AlNH₄(SO₄)₂, Al(OH)₃, AlPO₄, silica, kaolin; saponin derivative; carbon polynucleotides i.e. poly IC and poly AU and also detoxified cholera toxin (CTB) and E.coli heat
- labile toxin for induction of mucosal immunity. Preferred adjuvants include QuilATM, AlhydrogelTM and AdjuphosTM.

 Vaccines of the invention may be administered parenterally by injection, rapid infusion, nasopharyngeal absorption, dermoabsorption, or bucal or oral.

Vaccine compositions of the invention are used for the treatment or prophylaxis of streptococcus infection and/or diseases and symptoms mediated by streptococcus infection, in particular group A streptococcus (pyogenes), group B streptococcus (GBS or agalactiae), dysgalactiae, uberis, nocardia as well as Staphylococcus aureus. General information about Streptococcus is available in Manual of Clinical Microbiology by P.R.Murray et al. (1995, 6th Edition, 10 ASM Press, Washington, D.C.). More particularly group B streptococcus, agalactiae. In a particular embodiment vaccines are administered to those individuals at risk of GBS infection such as pregnant women and infants for sepsis, meningitis and pneumonia as well as immunocompromised 15 individuals such as those with diabetes, liver disease or cancer. Vaccines may also have veterinary applications such as for the treatment of mastitis in cattle which is mediated by the above mentioned bacteria as well as E.coli.

The vaccine of the present invention can also be used for the manufacture of a medicament used for the treatment or prophylaxis of streptococcus infection and/or diseases and symptoms mediated by streptococcus infection, in particular group A streptococcus (pyogenes), group B streptococcus (GBS or agalactiae), dysgalactiae, uberis, nocardia as well as Staphylococcus aureus. More particularly group B streptococcus, agalactiae.

Vaccine compositions are preferably in unit dosage form of about 0.001 to 100 µg/kg (antigen/body weight) and more preferably 0.01 to 10 µg/kg and most preferably 0.1 to 1 µg/kg 1 to 3 times with an interval of about 1 to 12 weeks intervals between immunizations, and more preferably 1 to 6

weeks.

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According to another aspect, there is provided polynucleotides encoding polypeptides of group B

- 5 streptococcus (GBS) characterized by the amino acid sequence selected from the group consisting of:
 - SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,
 - SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10,
 - SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15,
- 10 SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19,
 - SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24,
 - SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29,
 - SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34,
 - SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39,
- SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.

Preferred polynucleotides are those illustrated in figures la (SEQ ID NO: 1), 2a (SEQ ID NO: 7), 3a (SEQ ID NO: 13), 4a (SEQ ID NO: 22), 5a (SEQ ID NO: 27), 6a (SEQ ID NO: 32), 7a (SEQ ID NO: 37), 8 (SEQ ID NO: 42) and 9 (SEQ ID NO: 43) which correspond to the open reading frames, encoding polypeptides of the invention.

- Preferred polynucleotides are those illustrated in figures 1a (SEQ ID NO: 1), 2a (SEQ ID NO: 7), 3a (SEQ ID NO: 13), 4a (SEQ ID NO: 22), 5a (SEQ ID NO: 27), 6a (SEQ ID NO: 32), 7a (SEQ ID NO: 37), 8 (SEQ ID NO: 42) and 9(SEQ ID NO: 43) and fragments, analogues and derivatives thereof.
 - More preferred polynucleotides of the invention are those illustrated in Figures 7 (SEQ ID NO: 37), 8 (SEQ ID NO: 42) and 9 (SEQ ID NO: 43).
- Most preferred polynucleotides of the invention are those illustrated in Figures 8 (SEQ ID NO : 42) and 9 (SEQ ID NO :

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It will be appreciated that the polynucleotide sequences illustrated in the figures may be altered with degenerate codons yet still encode the polypeptides of the invention.

Due to the degeneracy of nucleotide coding sequences, other polynucleotide sequences which encode for substantially the same polypeptides of the present invention may be used in the practice of the present invention. These include but are not limited to nucleotide sequences which are altered by the substitution of different codons that encode the same amino acid residue within the sequence, thus producing a silent change.

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Accordingly the present invention further provides polynucleotides which hybridize to the polynucleotide sequences herein above described (or the complement sequences thereof) having 50% and preferably at least 70% identity between sequences. More preferably polynucleotides are hybridizable under stringent conditions i.e. having at least 95% identity and most preferably more than 97% identity.

By capable of hybridizing under stringent conditions is meant annealing of a nucleic acid molecule to at least a region of a second nucleic acid sequence (whether as cDNA, mRNA, or genomic DNA) or to its complementary strand under standard conditions, e.g. high temperature and/or low salt content, which tend to disfavor hybridization of noncomplementary nucleotide sequences. A suitable protocol is described in Maniatis T. et al., Molecular cloning: A Laboratory Manual, Cold Springs Harbor Laboratory, 1982, which is herein incorporated by reference.

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In a further aspect, polynucleotides encoding polypeptides

of the invention, or fragments, analogs or derivatives thereof, may be used in a DNA immunization method. That is, they can be incorporated into a vector which is replicable and expressible upon injection thereby producing the antigenic polypeptide in vivo. For example polynucleotides may be incorporated into a plasmid vector under the control of the CMV promoter which is functional in eukaryotic cells. Preferably the vector is injected intramuscularly.

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According to another aspect, there is provided a process for producing polypeptides of the invention by recombinant techniques by expressing a polynucleotide encoding said polypeptide in a host cell and recovering the expressed polypeptide product. Alternatively, the polypeptides can be produced according to established synthetic chemical techniques i.e. solution phase or solid phase synthesis of oligopeptides which are ligated to produce the full polypeptide (block ligation).

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For recombinant production, host cells are transfected with vectors which encode the polypeptide, and then cultured in a nutrient media modified as appropriate for activating promoters, selecting transformants or amplifying the genes. 25 Suitable vectors are those that are viable and replicable in the chosen host and include chromosomal, non-chromosomal and synthetic DNA sequences e.g. bacterial plasmids, phage DNA, baculovirus, yeast plasmids, vectors derived from combinations of plasmids and phage DNA. The polypeptide sequence may be incorporated in the vector at the 30 appropriate site using restriction enzymes such that it is operably linked to an expression control region comprising a promoter, ribosome binding site (consensus region or Shine-Dalgarno sequence), and optionally an operator (control element). One can select individual components of the 35 expression control region that are appropriate for a given

host and vector according to established molecular biology principles (Sambrook et al, Molecular Cloning: A Laboratory Manual, 2nd ed., Cold Spring Harbor, N.Y., 1989 incorporated herein by reference). Suitable promoters include but are not limited to LTR or SV40 promoter, E.coli lac, tac or trp promoters and the phage lambda P. promoter. Vectors will preferably incorporate an origin of replication as well as selection markers i.e. ampicillin resistance gene. Suitable bacterial vectors include pET, pQE70, pQE60, pQE-9, pbs, pD10 phagescript, psiX174, pbluescript SK, pbsks, pNH8A, 10 pNH16a, pNH18A, pNH46A, ptrc99a, pKK223-3, pKK233-3, pDR540, pRIT5 and eukaryotic vectors pBlueBacIII, pWLNEO, pSV2CAT, pOG44, pXT1, pSG, pSVK3, pBPV, pMSG and pSVL. Host cells may be bacterial i.e. E.coli, Bacillus subtilis, Streptomyces; fungal i.e. Aspergillus niger, Aspergillus 15 nidulins; yeast i.e. Saccharomyces or eukaryotic i.e. CHO,

Upon expression of the polypeptide in culture, cells are
typically harvested by centrifugation then disrupted by
physical or chemical means (if the expressed polypeptide is
not secreted into the media) and the resulting crude extract
retained to isolate the polypeptide of interest.
Purification of the polypeptide from culture media or lysate
may be achieved by established techniques depending on the
properties of the polypeptide i.e. using ammonium sulfate or
ethanol precipitation, acid extraction, anion or cation
exchange chromatography, phosphocellulose chromatography,
hydrophobic interaction chromatography, hydroxylapatite
chromatography and lectin chromatography. Final

COS.

The polypeptide may be expressed with or without a leader or secretion sequence. In the former case the leader may be removed using post-translational processing (see US

purification may be achieved using HPLC.

4,431,739; 4,425,437; and 4,338,397 incorporated herein by reference) or be chemically removed subsequent to purifying the expressed polypeptide.

- According to a further aspect, the GBS polypeptides of the invention may be used in a diagnostic test for streptococcus infection in particular GBS infection. Several diagnostic methods are possible, for example detecting streptococcus organism in a biological sample, the following procedure may be followed:
 - a) obtaining a biological sample from a patient;
 - b) incubating an antibody or fragment thereof reactive with a GBS polypeptide of the invention with the biological sample to form a mixture; and
- 15 c) detecting specifically bound antibody or bound fragment in the mixture which indicates the presence of streptococcus.

Alternatively, a method for the detection of antibody specific to a streptococcus antigen in a biological sample containing or suspected of containing said antibody may be performed as follows:

- a) isolating a biological sample from a patient;
- b) incubating one or more GBS polypeptides of the invention or fragments thereof with the biological sample to form a mixture; and

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c) detecting specifically bound antigen or bound fragment in the mixture which indicates the presence of antibody specific to streptococcus.

One of skill in the art will recognize that this diagnostic test may take several forms, including an immunological test such as an enzyme-linked immunosorbent assay (ELISA), a radioimmunoassay or a latex agglutination assay, essentially to determine whether antibodies specific for the protein are present in an organism.

The DNA sequences encoding polypeptides of the invention may also be used to design DNA probes for use in detecting the presence of streptococcus in a biological sample suspected of containing such bacteria. The detection method of this invention comprises:

- a) isolating the biological sample from a patient;
- b) incubating one or more DNA probes having a DNA sequence encoding a polypeptide of the invention or fragments thereof with the biological sample to form a mixture; and
- c) detecting specifically bound DNA probe in the mixture which indicates the presence of streptococcus bacteria.
- The DNA probes of this invention may also be used for detecting circulating streptococcus i.e. GBS nucleic acids in a sample, for example using a polymerase chain reaction, as a method of diagnosing streptococcus infections. The probe may be synthesized using conventional techniques and may be immobilized on a solid phase, or may be labeled with a detectable label. A preferred DNA probe for this application is an oligomer having a sequence complementary to at least about 6 contiguous nucleotides of the GBS polypeptides of the invention.

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Another diagnostic method for the detection of streptococcus in a patient comprises:

- a) labeling an antibody reactive with a polypeptide of the invention or fragment thereof with a detectable label;
- 30 b) administering the labeled antibody or labeled fragment to the patient; and
 - c) detecting specifically bound labeled antibody or labeled fragment in the patient which indicates the presence of streptococcus.

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A further aspect of the invention is the use of the GBS

polypeptides of the invention as immunogens for the production of specific antibodies for the diagnosis and in particular the treatment of streptococcus infection. Suitable antibodies may be determined using appropriate screening methods, for example by measuring the ability of a particular antibody to passively protect against streptococcus infection in a test model. One example of an animal model is the mouse model described in the examples herein. The antibody may be a whole antibody or an antigenbinding fragment thereof and may in general belong to any 10 immunoglobulin class. The antibody or fragment may be of animal origin, specifically of mammalian origin and more specifically of murine, rat or human origin. It may be a natural antibody or a fragment thereof, or if desired, a recombinant antibody or antibody fragment. The term 15 recombinant antibody or antibody fragment means antibody or antibody fragment which were produced using molecular biology techniques. The antibody or antibody fragments may be polyclonal, or preferably monoclonal. It may be specific for a number of epitopes associated with the GBS 20 polypeptides but is preferably specific for one.

EXAMPLE 1 Murine model of lethal Group B Streptococcus (GBS) infection

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The mouse model of GBS infection is described in detail in Lancefield et al (J Exp Med 142:165-179,1975). GBS strain C388/90 (Clinical isolate obtained in 1990 from the cephalorachidian fluid of a patient suffering from meningitis, Children's Hospital of Eastern Ontario, Ottawa, Canada) and NCS246 (National Center for Streptococcus, Provincial Laboratory of Public Health for Northern Alberta, Edmonton, Canada) were respectively serotyped as type Ia/c and type II/R.

To increase their virulence, the GBS strains C388/90 (serotype Ia/c) and NCS 246 (serotype II/R) were serially passaged through mice as described previously (Lancefield et al. J Exp Med 142:165-179, 1975). Briefly, the increase of virulence was monitored using intraperitoneal inoculations of serial dilutions of a subculture in Todd-Hewitt broth obtained from either the blood or spleen of infected mice. After the last passage, infected blood samples were used to inoculate Todd-Hewitt broth. After an incubation of 2 hours at 37°C with 7% CO2, glycerol at a final concentration of 10 10% (v/v) was added to the culture. The culture was then aliquoted and stored at -80° C for use in GBS challenge experiments. The number of cfu of GBS present in these frozen samples was determined. The bacterial concentration necessary to kill 100% (LD100) of the 18 weeks old mice were 15 determined to be 3.5X10⁵ and 1.1X10⁵ respectively for GBS strain C388/90 and NCS246, which corresponded to a significant increase in virulence for both strains. Indeed, the LD100 recorded before the passages for these two strains was higher than 10° cfu. 20

In a bacterial challenge, a freshly thawed aliquot of a virulent GBS strain was adjusted to the appropriate bacterial concentration using Todd-Hewitt broth and 1ml was 25 injected intraperitoneally to each female CD-1 mouse. The mice used for the passive protection experiments were 6 to 8 weeks old, while the ones used for the active protection experiments were approximately 18 weeks old at the time of the challenge. All inocula were verified by colony counts. Animals were observed for any sign of infection four times 30 daily for the first 48 h after challenge and then daily for the next 12 days. At the end of that period, blood samples were obtained from the survivors and frozen at -20°C. spleen obtained from each mouse that survived the challenge was cultured in order to identify any remaining GBS. 35

EXAMPLE 2 Immunization and protection in mice with formaldehyde killed whole GBS cells

- Formaldehyde killed GBS whole cells were prepared according to the procedures described in Lancefield et al (J Exp Med 142:165-179,1975). Briefly, an overnight culture on sheep blood agar plates (Quelab Laboratories, Montreal, Canada) of a GBS strain was washed twice in PBS buffer (phosphate buffered-saline, pH7.2), adjusted to approximately 3X10° cfu/mL and incubated overnight in PBS containing 0.3% (v/v) formaldehyde. The killed GBS suspension was washed with PBS and kept frozen at -80°C.
- 15 Female CD-1 mice, 6 to 8 weeks old (Charles River, St-Constant, Québec, Canada), were injected subcutaneously three times at two weeks interval with 0.1 ml of formaldehyde killed cells of GBS strain C388/90 (~6X10'GBS), or 0.1 ml of PBS for the control group. On the day before the immunization, AlhydrogelTM (Superfos Biosector, Frederikssund, Denmark) at a final concentration of 0.14 mg or 0.21 mg of Al, was added to these preparations and incubated overnight at 4°C with agitation. Serum samples were obtained from each mouse before the beginning of the

immunization protocol and two weeks after the last

injection. The sera were frozen at -20°C.

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Eight mice in each control group injected with PBS and the group immunized with formaldehyde killed whole cells GBS strain C388/90 (Ia/c) were challenged with 1.5X104 cfu of GBS strain C388/90 (Ia/c) one week after the third injection. All mice immunized with the formaldehyde killed GBS whole cells survived the homologous challenge while, within 5 days after the challenge, only 4 out of the 8 mice injected with PBS survived from the infection. In order to increase the mortality rate in the control groups, the

bacterial suspension had to be adjusted according to the age of the mice at the time of the bacterial challenge. In subsequent challenge experiments, when mice were older than 15 weeks, the bacterial inoculum was increased to concentrations between 3.0X10⁵ and 2.5X10⁶ cfu.

Table 1 Immunization of CD1 mice with formaldehyde killed whole cells of GBS and subsequent homologous challenge [strain C388/90 (Ia/c)] and heterologous challenge [strain NCS246 (II/R)].

antigenic preparations used for immunization ¹	number of living mice 14 days after the bacterial challenge (% Survival)					
	homologous challenge: strain C388/90 (la/c)	heterologous challenge: strain NCS246 (II/R)				
1st infection	st infection					
formaldehyde killed cells of GBS strain C388/90 (la/c) ²	8/8 (100) ³	n.d. ⁵				
control PBS	4/8 (50)	n.d.				
2nd infection	nd infection					
formaldehyde killed cells of GBS strain C388/90 (la/c)	6/6 (100) ⁴	0/6 (0) ⁶				
control PBS	2/6 (33)	0/6 (0)				

¹ alhydrogel™ at a final concentration of 0.14 mg or 0.21mg of Al was used;

² approximately 6X10⁷ cfu;

⁵ not done;

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In another experiment, one group of 12 mice corresponding to a control group was injected with PBS, while a second group of 12 mice was immunized with formaldehyde killed whole cells of GBS strain C388/90 (Ia/c). Six mice from each of these two groups were challenged with 2.1X10⁶ cfu of the GBS strain C388/90 (Ia/c) (Table I). As the first challenge experiment, all mice immunized with the GBS strain C388/90 (Ia/c) survived the homologous challenge. Only two out of the 6 mice injected with PBS survived the infection.

intraperitoneal challenge with 1 mL Todd-Hewitt culture medium containing GBS C388/90 (la/c) suspension adjusted to 1.5X10⁴ cfu;

⁴ intraperitoneal challenge with 1 mL Todd-Hewitt culture medium containing GBS C388/90 (la/c) suspension adjusted to 2.1X10⁶ cfu;

⁶ intraperitoneal challenge with 1 mL Todd-Hewitt culture medium containing GBS NCS246 (II/R) suspension adjusted to 1.2X10⁵ cfu.

The remaining 6 mice in both groups were then used one week later to verify whether this antigenic preparation could confer cross protection against strain NCS246 (II/R) which produce a serologically distinct capsule. None of the mice infected with this second GBS strain survived the infection. The later result suggested that most of the protective immune response induced by formaldehyde killed strain C388/90 is directed against the capsular polysaccharide and that it could be restricted to strains of that particular serotype. These results clearly indicated that this particular model of infection can be efficiently used to study the protection conferred by vaccination.

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EXAMPLE 3 Immunization of rabbit with formaldehyde killed whole GBS cells and passive protection in mice

A New Zealand rabbit (2.5 kg, Charles River, St-Constant, Québec, Canada) was immunized with formaldehyde killed 20 cells of GBS strain C388/90 (Ia/c) to obtain hyperimmune This rabbit was injected subcutaneously three serum. times at three weeks interval with approximately 1.5X109 cfu of formaldehyde killed whole cells of GBS strain C388/90 (Ia/c). Freund's complete adjuvant (Gibco BRL 25 Life Technologies, Grand Island, New York) was used as the adjuvant for the first immunization, while Freund's incomplete adjuvant (Gibco BRL) was used for the following two injections. Serum samples were obtained before the beginning of the immunization protocol and two weeks after 30 the last injection. The sera were frozen at -20°C.

The ability of this particular rabbit hyperimmune serum to passively protect mice against a lethal infection with GBS

was also evaluated. Intraperitoneal injection of mice with either 15 or 25 μ L of hyperimmune rabbit serum 18 hours before the challenge protected 4 out of 5 mice (80%) against the infection. Comparatively, survival rates lower than 20% were recorded for mice in the control group injected with PBS or serum obtained from a rabbit immunized with meningococcal outer membrane preparation. This result clearly indicates that the immunization of another animal species with killed GBS cells can induce the production of antibodies that can passively protect mice. This reagent will also be used to characterize clones.

Table 2 Passive protection of CD-1 mice conferred by rabbit serum obtained after immunization with formaldehyde killed group B whole streptococci (strain C388/90 (Ia/c)) antigenic preparation

groups	number of living mice 14 days after the bacterial challenge with GBS strain C388/90 (Ia/c) ²	% survival
rabbit hyperimmune serum² - 25 μl	4/5	80
rabbit hyperimmune serum¹ - 15 μl	4/5	80
control rabbit serum - 25 μl	1/5	20
control PBS	1/10	10

Freund's complete adjuvant was used for first immunization, and Freund's incomplete adjuvant for the following two injections;

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intraperitoneal challenge with 1 ml Todd-Hewitt culture medium containing GBS C388/90 (Ia/c) suspension adjusted to 2X104 cfu.

EXAMPLE 4 Recombinant production of His.Tag-GBS fusion protein

The coding region of a GBS gene was amplified by PCR (DNA Thermal Cycler GeneAmp PCR system 2400 Perkin Elmer, San Jose, CA) from the genomic DNA of GBS strain C388/90 (Ia/c) using the oligos that contained base extensions for the addition of the restriction sites BglII (AGATCT) and HindIII (AAGCTT), respectively. The PCR product was purified from agarose gel using a Qiaex II gel extraction kit from Qiagen 10 (Chatsworth, CA), digested with the restriction enzymes BglII and HindIII (Pharmacia Canada Inc Baie d'Urfe, Canada), and extracted with phenol:chloroform before ethanol precipitation. The pET-32b(+) vector (Novagen, Madison, WI) containing the thioredoxin-His. Tag sequence was digested 15 with the restriction enzymes BglII and HindIII, extracted with phenol:chloroform, and then ethanol precipitated. The BglII-HindIII genomic DNA fragment was ligated to the BglII-HindIII pET-32b(+) vector to create the coding sequence for thioredoxin-His.Tag-GBS fusion protein whose gene was under 20 control of the T7 promoter. The ligated products were transformed into E. coli strain XLI Blue MRF' (\Delta(mcrA) 183\Delta (mcrCB-hsdSMR-mrr)173 endAl supE44 thi-1 recAl gyrA96 relAl lac [F'proAB lacIqZΔM15Tn10 (Tetr)]c) (Stratagene, La Jolla, CA) according to the method of Simanis (Hanahan, D. DNA 25 Cloning, 1985, D.M. Glover (ed.), pp. 109-135). The recombinant pET plasmid was purified using a Qiagen kit (Qiagen, Chatsworth, CA) and the nucleotide sequence of the DNA insert was verified by DNA sequencing (Taq Dye Deoxy Terminator Cycle Sequencing kit, ABI, Foster City, CA). The 30 recombinant pET plasmid was transformed by electroporation (Gene Pulser II apparatus, BIO-RAD Labs, Mississauga, Canada) into E. coli strain AD494 (DE3) (∆ara leu7697 ΔlacX74 ΔphoA PvuII phoR ΔmalF3 F'[lac*(lacIq) pro] 35 trxB::Kan (DE3)) (Novagen, Madison, WI). In this strain of

E. coli, the T7 promoter controlling expression of the fusion protein, is specifically recognized by the T7 RNA polymerase (present on the $\lambda DE3$ prophage) whose gene is under the control of the lac promoter which is inducible by isopropyl- β -D-thio-galactopyranoside (IPTG).

The transformant AD494(DE3)/rpET was grown at 37°C with agitation at 250 rpm in LB broth (peptone 10g/L, Yeast extract 5g/L, NaCl 10g/L) containing 100µg of ampicillin (Sigma-Aldrich Canada Ltd., Oakville, Canada) per mL until the A₆₀₀ reached a value of 0.6. In order to induce the production of the thioredoxin-His.Tag-GBS fusion protein, the cells were incubated for 2 additional hours in the presence of IPTG at a final concentration of 1mM. The bacterial cells were harvested by centrifugation.

The recombinant fusion protein produced by AD494(DE3)/rpET32 upon IPTG induction for 2h was partially obtained as insoluble inclusion bodies which were purified from endogenous E. coli proteins by the isolation of insoluble 20 aggregates (Gerlach, G.F. et al 1992, Infect. Immun. 60:892). Induced cells from a 500 mL culture were resuspended in 20 mL of 25% sucrose-50mM Tris-HCl buffer (pH8.0) and frozen at -70°C. Lysis of cells in thawed suspension was achieved by the addition of 5mL of a solution 25 of lysozyme (10mg/mL) in 250mM Tris-HCl buffer (pH8.0) followed by an incubation of 10 to 15 min on ice, and the addition of 150mL of detergent mix (5 parts of 20mM Tris-HCl buffer [pH7.4]-300mM NaCl-2% deoxycholic acid-2% Nonidet P-40 and 4 parts of 100mM Tris-HCl buffer [pH8]-50mM EDTA-2% 30 Triton X-100) followed by 5 min incubation on ice. Upon sonication, protein aggregates were harvested by centrifugation for 30 min at 35,000 X g and a sample of the soluble cellular fraction was kept. The aggregated proteins were solubilized in 6M guanidine hydrochloride. 35

presence of the fusion protein in both the soluble and insoluble fractions was shown by Western Blot analysis using the serum of a mouse injected with formaldehyde killed cells of GBS strain C388/90 (Ia/c) that survived a bacterial challenge with the corresponding GBS strain.

The purification of the fusion protein from the soluble fraction of IPTG-induced AD494 (DE3) / rpET was done by affinity chromatography based on the properties of the 10 His. Tag sequence (6 consecutive histidine residues) to bind to divalent cations (Ni2+) immobilized on the His.Bind metal chelation resin (Novagen, Madison, WI). The purification method used are those described in the pET system Manual, 6th Edition (Novagen, Madison, WI). Briefly, the pelleted cells obtained from a 100mL culture induced with IPTG was 15 resuspended in 4mL of Binding buffer (5mM imidazole-500mM NaCl-20mM Tris-HCl pH7.9), sonicated, and spun at 39,000 X g for 20 min to remove debris. The supernatant was filtered $(0.45\mu m \text{ pore size membrane})$ and deposited on a column of His.Bind resin equilibrated in Binding buffer. The column 20 was then washed with 10 column volumes of Binding buffer followed by 6 column volumes of Wash buffer (20mM imidazole-500mM NaCl-20mM Tris-HCl pH7.9). The thioredoxin-His.Tag-GBS fusion protein was eluted with Elute buffer (1M 25 imidazole-500mM NaCl-20mM Tris-HCl pH7.9). The removal of the salt and imidazole from the sample was done by dialysis against 3 X 1 liter PBS at 4°C.

The quantities of fusion protein obtained from either the soluble or insoluble cytoplasmic fractions of *E. coli* were estimated by Coomassie staining of a sodium dodecyl sulfate (SDS)-polyacrylamide gel with serial dilutions of these proteins and a bovine serum albumin standard (Pierce Chemical Co. Rockford, Ill.).

EXAMPLE 5 Recombinant production of GBS protein under control of lambda P_L promoter

The DNA coding region of a GBS protein was inserted
downstream of the promoter λP_L into the translation vector pURV22. This plasmid was derived from p629 (George et al, 1987, Bio/Technology 5:600) from which the coding region for a portion of the herpes simplex virus type I (HSV-I) glycoprotein (gD-1) was removed and the ampicillin
resistance gene replaced by a kanamycin cassette obtained from the plasmid vector pUC4K (Pharmacia Biotech Canada Inc., Baie D'Urfe, Canada). The vector contained a cassette of the bacteriophage λ cI857 temperature sensitive repressor gene from which the functional P_R promoter had been deleted.
The inactivation of the cI857 repressor by temperature increase from the ranges of 30-37°C to 37-42°C resulted in

increase from the ranges of 30-37°C to 37-42°C resulted in the induction of the gene under the control of λ P_L. The translation of the gene was controlled by the ribosome binding site cro followed downstream by a BglII restriction site (AGATCT) and the ATG: ACTAAGGAGGTTAGATCTATG.

Restriction enzymes and T4 DNA ligase were used according to suppliers (Pharmacia Biotech Canada Inc., Baie D'Urfe, Canada; and New England Biolabs Ltd., Mississauga, Canada).

- Agarose gel electrophoresis of DNA fragments was performed as described by Sambrook et al. (Molecular cloning : A laboratory Manual, 1989, Cold Spring Harbor Laboratory Press, N.Y). Chromosomal DNA of the GBS bacteria was prepared according to procedures described in Jayarao et al
- (J. Clin. Microbiol., 1991, 29:2774). DNA amplification reactions by polymerase chain reaction (PCR) were made using DNA Thermal Cycler GeneAmp PCR system 2400 (Perkin Elmer, San Jose, CA). Plasmids used for DNA sequencing were purified using plasmid kits from Qiagen (Chatsworth, CA).
- 35 DNA fragments were purified from agarose gels using Qiaex II

gel extraction kits from Qiagen (Chatsworth, CA). Plasmid transformations were carried out by the method described by Hanahan (DNA Cloning, Glover (ed.) pp, 109-135, 1985). The sequencing of genomic DNA inserts in plasmids was done using synthetic oligonucleotides which were synthesized by 5 oligonucleotide synthesizer model 394 (the Perkin-Elmer Corp., Applied Biosystems Div. (ABI), Foster City, CA). The sequencing reactions were carried out by PCR using the Taq Dye Deoxy Terminator Cycle Sequencing kit (ABI, Foster City, 10 CA) and DNA electrophoresis was performed on automated DNA sequencer 373A (ABI, Foster City, CA). The assembly of the DNA sequence was performed using the program Sequencer 3.0 (Gene Codes Corporation, Ann Arbor, MI). Analysis of the DNA sequences and their predicted polypeptides was performed with the program Gene Works version 2.45 (Intelligenetics, 15 Inc., Mountain View CA).

The coding region of the GBS gene was amplified by PCR from GBS strain C388/90 (Ia/c) genomic DNA using oligos that contained base extensions for the addition of restriction 20 sites BglII (AGATCT) and XbaI(TCTAGA), respectively. The PCR product was purified from agarose gel using a Qiaex II gel extraction kit from Qiagen (Chatsworth, CA), digested with the restriction enzymes BglII and XbaI, and extracted with phenol:chloroform before ethanol precipitation. The pURV22 25 vector was digested with the restriction enzymes BglII and XbaI, extracted with phenol:chloroform, and ethanol precipitated. The BglII-XbaI genomic DNA fragment was ligated to the BglII-XbaI pURV22 vector in which the GBS gene was under the control of the λPL promoter. The ligated 30 products were transformed into E. coli strain XLI Blue MRF' $(\Delta (mcrA) 183\Delta (mcrCB-hsdSMR-mrr) 173 endA1 supE44 thi-1 recA1$ gyrA96 relA1 lac[F' proAB lac1qZAM15 Tn10(Tetr)]c) (Stratagene, La Jolla CA) according to the methods described in Hanahan, supra. Transformants harboring plasmids with the 35

insert were identified by analysis of lysed cells submitted to electrophoresis on agarose gel (Sambrook et al, <u>supra</u>). The recombinant pURV22 plasmid was purified using a Qiagen kit (Qiagen, Chatsworth, CA) and the nucleotide sequence of the DNA insert was verified by DNA sequencing.

The transformant XLI Blue MRF'/rpURV22 was grown at 34°C with agitation at 250 rpm in LB broth containing $50\mu g$ of kanamycin per mL until the A_{600} reached a value of 0.6. In order to induce the production of the fusion protein, the cells were incubated for 4 additional hours at 39°C. The bacterial cells were harvested by centrifugation , resuspended in sample buffer, boiled for 10 min and kept at -20°C.

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EXAMPLE 6 Subcloning GBS protein gene in CMV plasmid pCMV-GH

The DNA coding region of a GBS protein was inserted in phase downstream of the human growth hormone (hGH) gene which was under the transcriptional control of the cytomegalovirus (CMV) promoter in the plasmid vector pCMV-GH (Tang et al, Nature, 1992, 356:152). The CMV promoter is non functional in E. coli cells but active upon administration of the plasmid in eukaryotic cells. The vector also incorporated the ampicillin resistance gene.

The coding region of the gene was amplified by PCR from genomic DNA of GBS strain C388/90 (Ia/c) using the oligos that contained base extensions for the addition of the restriction sites BglII (AGATCT) and HindIII (AAGCTT). The PCR product was purified from agarose gel using a Qiaex II gel extraction kit from Qiagen (Chatsworth, CA), digested with the restriction enzymes BglII and HindIII, and extracted with phenol:chloroform before ethanol precipitation. The pCMV-GH vector (Laboratory of Dr. Stephen

A. Johnston, Department of Biochemistry, The University of Texas, Dallas, Texas) containing the human growth hormone to create fusion proteins was digested with the restriction enzymes BamHI and HindIII, extracted with phenol:chloroform, and ethanol precipitated. The 1.3-kb BglII-HindIII genomic DNA fragment was ligated to the BamHI -HindIII pCMV-GH vector to create the hGH-GBS fusion protein under the control of the CMV promoter. The ligated products were transformed into E. coli strain DH5 α [ϕ 80 lacZ Δ M15 endA1 recAl hsdR17 ("K-"K+) supE44 thi-1λ gyrA96 relAl Δ(lacZYA-10 argF) U169] (Gibco BRL, Gaithersburg, MD) according to the methods described by Hanahan, supra. Transformants harboring plasmids with the insert were identified by analysis of lysed cells submitted to electrophoresis on agarose gel (Sambrook, J. et al , supra). The recombinant 15 pCMV plasmid was purified using a Qiagen kit (Qiagen, Chatsworth, CA) and the nucleotide sequence of the DNA insert was verified by DNA sequencing.

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EXAMPLE 7 Immunological activity of GBS protein to GBS challenge

Four groups of 12 female CD-1 mice (Charles River, St25 Constant, Quebec, Canada) of 6 to 8 weeks were injected subcutaneously three times at three week intervals with 0.1mL of the following antigenic preparations: formaldehyde killed cells of GBS strain C388/90 (~6X10⁷ cfu), 20μg of thioredoxin-His.Tag-GBS fusion protein obtained from the insoluble (inclusion bodies) or 20μg of the fusion protein, affinity purified (nickel column), from the soluble cytoplasmic fraction in E.coli, or 20μg of affinity purified (nickel column) thioredoxin-His.Tag control polypeptide. 20μg of QuilATM (Cedarlane Laboratories Ltd, Hornby, Canada)

was added to each antigenic preparation as the adjuvant. Serum samples were obtained from each mouse before immunization (PB) and on days 20 (TB1), 41 (TB2) and 54 (TB3) during the immunization protocols. Sera were frozen at -20°C.

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An increase of the ELISA titers was recorded after each injection of the fusion protein indicating a good primary response and a boost of the specific humoral immune response after each of the second and third administration. At the 10 end of the immunization period, the means of reciprocal ELISA titers was 456,145 for the group immunized with 20µg of fusion protein obtained from inclusion bodies compared to 290,133 for the group of mice immunized with the protein from soluble fraction in E.coli. The latter result suggests 15 that the protein obtained from inclusion bodies could be more immunogenic than the soluble protein. Analysis of mice sera in ELISA using the affinity purified thioredoxin-His. Tag to coat plates showed that negligible antibody titers are made against the thioredoxin-His. Tag portion of 20 the fusion protein. The reactivity of the sera from mice injected with the recombinant fusion protein was also tested by ELISA against formaldehyde killed whole cells of GBS strain C388/90. The antibodies induced by immunization with recombinant fusion protein also recognized their specific 25 epitopes on GBS cells indicating that their conformation is close enough to the native streptococcal protein to induce cross-reactive antibodies.

To verify whether the immune response induced by immunization could protect against GBS infection, mice were challenged with 3.5X10⁵ cfu of GBS strains C338/90(Ia/c) and 1.2X10⁵ cfu of strain NCS246(II/R) the results of which are illustrated in tables 3 and 4 respectively. Mice immunized with control thioredoxin-His.Tag peptide were not protected against challenge with either GBS strain while those

immunized with formaldehyde killed C388/90 whole cells only provided protection against homologous challenge. The thioredoxin-His.Tag-GBS fusion protein of the invention protected mice from challenge with both GBS strains. Blood and spleen culture of these mice did not reveal the presence of any GBS.

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Table 3 Survival from GBS strain C388/90 (Ia/c) challenge¹

immunizing agent	no. mice surviving challenge	% survival
thioredoxin-His.Tag²	1 / 6	17
formaldehyde killed C388/90 cells³	6 / 6	100
thioredoxin-His.Tag-GBS fusion (inclusion body preparation)4	6 / 6	100
thioredoxin-His.Tag-GBS fusion (cytoplasmic fraction)4	6 / 6	100

intraperitoneal administration with 1 ml Todd-Hewitt culture medium adjusted to 3.5X10⁵ cfu;

² 20µg administered; posterior legs paralyzed in surviving mouse; GBS detected in blood and spleen; 6X107 cfu administered;

^{4 20}μg administered.

Table 4 Survival from GBS strain NCS246 (II/R) challenge¹

immunizing agent	no. mice surviving challenge	% survival
thioredoxin-His.Tag2	0 / 6	0
formaldehyde killed C388/90 cells ³	2 / 6	34
thioredoxin-His.Tag-GBS fusion (inclusion body preparation) ²	5 / 5 ⁴	100
thioredoxin-His.Tag-GBS fusion (cytoplasmic fraction)2	6 / 6	100

intraperitoneal administration with 1 ml Todd-Hewitt culture medium containing GBS NCS246(II/R) suspension adjusted to 1.2X10⁵ cfu.

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EXAMPLE 8 Immunization with recombinant GBS protein confers protection against experimental GBS infection

This example illustrates the protection of mice against fatal GBS infection by immunization with the recombinant protein corresponding to the SEQ ID NO:39.

Groups of 10 female CD-1 mice (Charles River) were immunized subcutaneously three times at three-week intervals with 20 μg of recombinant protein purified from E. coli strain BLR (Novagen) harboring the recombinant pURV22 plasmid vector containing the GBS gene corresponding to SEQ ID NO:42 in presence of 20 μg of QuilATM adjuvant (Cedarlane Laboratories Ltd, Hornby, Canada) or, as control, with

² 20µg administered;

³ 6X10⁷ cfu administered;

^{10 4} one mouse died during immunization.

QuilATM adjuvant alone in PBS. Blood samples were collected from the orbital sinus on day 1, 22 and 43 prior to each immunization and fourteen days (day 57) following the third injection. One week later the mice were challenged with approximately 10⁴ to 10⁶ CFU of various virulent GBS strains. Samples of the GBS challenge inoculum were plated on TSA/5% sheep blood agar plates to determine the CFU and to verify the challenge dose. Deaths were recorded for a period of 14 days and on day 14 post-challenge, the surviving mice were sacrificed and blood and spleen were tested for the presence of GBS organisms. The survival data are shown in table 5.

Prechallenge sera were analyzed for the presence of antibodies reactive with GBS by standard immunoassays. Elisa and immunoblot analyses indicated that immunization with recombinant GBS protein produced in *E. coli* elicited antibodies reactive with both, recombinant and native GBS protein. Antibody responses to GBS are described in Example 9.

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Table 5. Ability of recombinant GBS protein corresponding to SEQ ID NO: 39 to elicit protection against 8 diverse GBS challenge strains

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	Challenge	strain		
Immunogen	Designation	Type	No. alive:	No. dead 1
rGBS protein none	C388/90	Ia/c	8 : 2 0 : 10	(P<0.0001)
rGBS protein none	NCS 246	II/R	10 : 0 3 : 7	(P=0.0012)
rGBS protein none	ATCC12401	Ib	10 : 0 3 : 7	(P=0.001)
rGBS protein none	NCS 535	V	10 : 0 5 : 5	(P=0.01)
rGBS protein none	NCS 9842	VI	10 : 0 0 : 10	(P<0.0001)
rGBS protein NCS 915-F ³ none	NCS 915	III	7 : 3 1 : 9 4 : 6	(P=0.0007) ²
rGBS protein NCS 954-F	NCS 954	III/R	7 : 3 4 : 6	(P=0.002)
none	•		1:9	
rGBS protein COH1-F	COH1	III	4 : 6 3 : 7	(P=0.0004)
none	·		0:10	

Groups of 10 mice per group were used, the number of mice surviving to infection and the number of dead mice are indicated. The survival curves corresponding to recombinant GBS protein-immunized animals were compared to the survival curves corresponding to mock-immunized animals using the log-rank test for nonparametric analysis.

All hemocultures from surviving mice were negative at day 14 20 post-challenge. Spleen cultures from surviving mice were negative except for few mice from experiment MB-11.

² Comparison analysis to NCS915-F-immunized animals.

^{15 &}lt;sup>3</sup> Animals were immunized with formaldehyde-killed GBS in presence of QuilATM adjuvant.

EXAMPLE 9 Vaccination with the recombinant GBS protein elicits an immune response to GBS

Groups of 10 female CD-1 mice were immunized subcutaneously with recombinant GBS protein corresponding to SEQ ID NO:39 as described in Example 8. In order to assess the antibody response to native GBS protein, sera from blood samples collected prior each immunization and fourteen days after the third immunization were tested for antibody reactive with GBS cells by ELISA using plates coated with 10 formaldehyde-killed GBS cells from type III strain NCS 954, type Ib strain ATCC12401, type V strain NCS 535 or type VI strain NCS 9842. The specificity of the raised antibodies for GBS protein was confirmed by Western blot analyses to GBS cell extracts and purified recombinant antigens. The 15 results shown in Figure 10 clearly demonstrate that animals respond strongly to recombinant GBS protein used as immunogens with median reciprocal antibody titers varying between 12000 and 128000, for sera collected after the third immunization, depending of the coating antigen. All 20 preimmune sera were negative when tested at a dilution of 1 :100. GBS-reactive antibodies were detectable in the sera of each animal after a single injection of recombinant GBS protein.

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Example 10 Antigenic conservation of the GBS protein of the present invention

Monoclonal antibodies (MAbs) specific to the GBS protein of the present invention were used to demonstrate that this surface antigen is produced by all GBS and that it is also antigenically highly conserved.

A collection of 68 GBS isolates was used to evaluate the reactivity of the GBS-specific MAbs. These strains were 10 obtained from the National Center for Streptococcus, Provincial Laboratory of Public Health for Northern Alberta, Canada; Centre Hospitalier Universitaire de Quebec, Pavillon CHUL, Quebec, Canada; American Type Culture Collection, USA; Laboratoire de Sante Publique du Quebec, Canada; and Dept. 15 of Infectious Disease, Children's Hospital and Medical Center, Seattle, USA. All eight Mabs were tested against the following panel of strains: 6 isolates of serotype Ia or Ia/c, 3 isolates of serotype Ib, 4 isolates of serotype II, 14 isolates of serotype III, 2 isolates of serotype IV, 2 20 isolates of serotype V, 2 isolates of serotype VI, 2 isolates of serotype VII, 1 isolate of serotype VIII, 10 isolates that were not serotyped and 3 bovine S. agalactiae strains. MAb 3A2 was also reacted with additional GBS: 9 isolates of serotype Ia/c and 10 isolates of serotype V. 25 The strains were grown overnight on blood agar plates at 37°C in an atmosphere of 5% CO2. Cultures were stored at -70°C in heart infusion broth with 20% (v/v) glycerol.

To obtain the GBS protein-specific MAbs, mice were immunized three times at three-week intervals with 20 μg of purified recombinant GBS protein (SEQ ID NO :44) in the presence of 20% QuilATM adjuvant. Hybridoma cell lines were generated by fusion of spleen cells recovered from immunized mice with the nonsecreting SP2/O myeloma cell line as described

previously (Hamel, J. et al. 1987. J. Med. Microbiol. 23:163-170). Hybrid clone supernatants were tested for specific antibody production by ELISA using formaldehyde inactivated GBS and purified recombinant GBS protein (SEQ ID NO :39 or 44) as coating antigen, as previously described (Hamel, J. et al. 1987. J. Med. Microbiol. 23:163-170). Specific hybrid were cloned by limiting dilutions, expanded, and frozen in liquid nitrogen. Production of recombinant GBS protein was presented in Examples 4 & 5. Purified recombinant GBS protein or formaldehyde inactivated GBS were 10 resolved by electrophoresis by using the discontinuous buffer system of Laemmli as recommended by the manufacturer and then transfer onto nitrocellulose membrane for Western immunoblotting as described previously (Martin et al. 1992. Infect. Immun. 60:2718-2725). 15

Western immunoblotting experiments clearly indicated that all eight MAbs recognized a protein band that corresponded to the purified recombinant GBS protein (SEQ ID NO :39).

20 These MAbs also reacted with a protein band present in every GBS isolates tested so far. The reactivity of these GBS-specific MAbs are presented in Table 6. Each MAb reacted well with all 46 GBS. In addition, these MAbs also recognized the 3 S. agalactiae strains of bovine origin that were tested. MAb 3A2 also recognized nineteen GBS; 9 isolates of serotype Ia/c and 10 of serotype V. The other MAbs were not tested against these additional strains.

These results demonstrated that the GBS protein (SEQ ID NO :39) was produced by all the 65 GBS and the three 3 S. agalactiae strains of bovine origin that were tested so far. More importantly, these results clearly demonstrated that the epitopes recognized by these eight GBS-specific MAbs were widely distributed and conserved among GBS. These results also indicated that these epitopes were not

restricted to serologically related isolates since representatives of all known GBS serotypes including the major disease causing groups were tested.

In conclusion, the data presented in this example clearly demonstrated that the GBS protein of the present invention is produced by all GBS and that it is antigenically highly conserved.

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agalactiae S_{\cdot} different MAbs with eight GBS protein-specific as evaluated by Western immunoblots Reactivity of . 9 Table

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- 0	rains	VIII	(1)		ŧ	-		-	-				•		,	-4
	riae st	VIT	(2)	2	3	2	C	7	2	2	7	0	3	7	c	7
	agalaci	VT	(2)	2	1	2		7	2	3	7	2	7	7	c	7
	ω.	Λ	(2)	í	4	2		~	2	7	~	C	7	7	C	~
	otype of	111	(2)	ì	1	2	3	2	C	7	2	C	7	2	(``
	ser	111	(7)	1	1 '	4	4	4		‡	4		4	4	'	7
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by recognized were serotype of strains 10 and Ia/c serotype 1 Nine additional strains of serot: MAb 3A2. 2 These strains were not serotyped

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WE CLAIM:

1. An isolated polynucleotide encoding a polypeptide having at least 70% identity to a second polypeptide having a sequence selected from the group consisting of:

```
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.
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- 2. A polynucleotide according to claim 1, wherein said polynucleotide encodes a polypeptide having at least 95% identity to the second polypeptide.
- An isolated polynucleotide encoding a polypeptide capable of generating antibodies having binding specificity for a polypeptide having a sequence selected from the group consisting of:

 SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.

4. An isolated polynucleotide that is complementary to the polynucleotide of claim 1.

- 5. An isolated polynucleotide that is complementary to the polynucleotide of claim 3.
- 6. The polynucleotide of claim 1, wherein said polynucleotide is DNA.
- 7. The polynucleotide of claim 3, wherein said polynucleotide is DNA.
- 8. The polynucleotide of claim 1, wherein said polynucleotide is RNA.
- 9. The polynucleotide of claim 3, wherein said polynucleotide is RNA.
- 10. A polynucleotide which hybridizes under stringent conditions to a second polynucleotide having a sequence selected from the group consisting of:

 SEQ ID NO: 1, SEQ ID NO: 7, SEQ ID NO: 13, SEQ ID NO: 22, SEQ ID NO: 27, SEQ ID NO: 32, SEQ ID NO: 37, SEQ ID NO: 42 and SEQ ID NO: 43 or fragments, analogues or derivatives thereof.
- 11. A polynucleotide which hybridizes under stringent conditions to a second polynucleotide having a sequence selected from the group consisting of : SEQ ID NO : 37, SEQ ID NO : 42 and SEQ ID NO : 43.
- 12. A polynucleotide according to claim 11 which hybridizes under stringent conditions to a second polynucleotide having the sequence SEQ ID NO : 37.

13. A polynucleotide according to claim 11 which hybridizes under stringent conditions to a second polynucleotide having the sequence SEQ ID NO : 42.

- 14. A polynucleotide according to claim 11 which hybridizes under stringent conditions to a second polynucleotide having the sequence SEQ ID NO : 43.
- 15. A polynucleotide according to claim 10 wherein said polynucleotide has at least 95% complementarity to the second polynucleotide.
- 16. A polynucleotide according to claim 11 wherein said polynucleotide has at least 95% complementarity to the second polynucleotide.
- 17. A vector comprising the polynucleotide of claim 1, wherein said polynucleotide is operably linked to an expression control region.
- 18. A vector comprising the polynucleotide of claim 3, wherein said polynucleotide is operably linked to an expression control region.
- 19. A host cell transfected with the vector of claim 17.
- 20. A host cell transfected with the vector of claim 18.
- 21. A process for producing a polypeptide comprising culturing a host cell according to claim 19 under conditions suitable for expression of said polypeptide.
- 22. A process for producing a polypeptide comprising culturing a host cell according to claim 20 under condition suitable for expression of said polypeptide.

23. An isolated polypeptide having at least 70% identity to a second polypeptide having a sequence selected from the group consisting of:

```
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:35, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.
```

- 24. The isolated polypeptide of claim 23 having a sequence according to SEQ ID NO: 39.
- 25. The isolated polypeptide of claim 23 having a sequence according to SEQ ID NO: 44.
- 26. An isolated polypeptide capable of generating antibodies having binding specificity for a second polypeptide having a sequence selected from the group consisting of:

```
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.
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27. The isolated polypeptide of claim 26 having a sequence according to SEQ ID NO: 39.

- 28. The isolated polypeptide of claim 26 having a sequence according to SEQ ID NO: 44.
- 29. An isolated polypeptide having an amino acid sequence selected from the group consisting of:

 SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39,
- 30. The isolated polypeptide of claim 29 having an amino acid sequence according to SEQ ID NO: 39.

derivatives thereof.

SEQ ID NO:40 and SEQ ID NO:41 or fragments, analogs or

- 31. An isolated polypeptide having an amino acid sequence according to SEQ ID NO: 44.
- 32. An isolated polypeptide according to any one of claims 29 to 31, wherein the N-terminal Met residue is deleted.
- 33. An isolated polypeptide according to any one of claims 29 to 30, wherein the secretory amino acid sequence is deleted.
- 34. A vaccine composition comprising a polypeptide according to any one of claims 23 to 31 and a pharmaceutically acceptable carrier, diluent or adjuvant.

35. A vaccine composition comprising a polypeptide according to claim 32 and a pharmaceutically acceptable carrier, diluent or adjuvant.

- 36. A vaccine composition comprising a polypeptide according to claim 33 and a pharmaceutically acceptable carrier, diluent or adjuvant.
- 37. A method for therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of a composition according to claim 34.
- 38. A method for therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of a composition according to claim 35.
- 39. A method for therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of a composition according to claim 36.
- 40. A method according to any one of claims 37 to 39, wherein said animal is a bovine.
- 41. A method according to any one of claims 37 to 39, wherein said animal is a human.

42. A method according to any one of claims 37 to 39, wherein said bacterial infection is selected from the group consisting of group A streptococcus and group B streptococcus.

- 43. A method according to claim 42, wherein said bacterial infection is group B streptococcus.
- 44. Use of a vaccine composition according to claim 34 for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to or infected with streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.
- 45. Use of a vaccine composition according to any one of claims 35 to 36 for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to or infected with streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.
- 46. Use of a vaccine composition according to any one claims 23 to 31 for the manufacture of a vaccine for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.
- 47. Use of a vaccine composition according to claim 32 for the manufacture of a vaccine for the therapeutic or

prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.

48. Use of a vaccine composition according to claim 33 for the manufacture of a vaccine for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.

S G K	E P A	N R F S	W A K	N K L	L I N G	60
•	AACTCTAGCA T L A	GCAACTATCT A T I L	TATTTTTTGC F F A	AGTTCAATTC V Q F	ATAGGTCTTA I G L K	120
AACCAGATTA P D Y	CCCTGGAAAA P G K	ACCTACTTTA T Y F I	TTATCCTATT I L L	GACAGCATGG T A W	ACTTTGATGG T L M A	180
CATTAGTAAC L V T	TGCTTTAGTG A L V	GGATGGGATA G W D N	ATAGGTATGG R Y G	TTCCTTCTTG S F L	TCGTTATTAA S L L I	240
TATTATTATT L L F	CCAGCTTGGT Q L G	TCAAGCGCAG S S A G	GAACTTACCC T Y P	AATAGAATTG I E L	AGTCCTAAGT S P K F	300
TCTTTCAAAC F Q T	AATTCAACCA I Q P	TTTTTACCGA F L P M	TGACTTACTC T Y S	TGTTTCAGGA V S G	TTAAGAGAGA L R E T	360
CCATCTCGTT I S L	GACGGGAGAC T G D	GTTAACCATC V N H Q	AATGGAGAAT W R M	GCTAGTAATC L V I	TTTTTAGTAT F L V S	420
CATCGATGAT S M I	ACTTGCTCTT L A L	CTTATTTATC L I Y R	GTAAACAAGA K Q E	AGATTAATAG D	AAAGTATCTA	480
GTGATAGACT	AACAGTATGA	TATGGTATGT	CAAAGTATTT	AGGAGGAGAA	GATATGTCTA M S T	540
CTTTAACAAT L T I	AATTATTGCA I I A	ACATTAACTG T L T A		TTTTTATATT F Y I	ATGTATTTGG M Y L E	600
AGACGTTAGC T L A	CACCCAGTCA T Q S	AATATGACTG N M T G	GGAAGATTTT K I F	TAGTATGTCT S M S	AAAGAAGAGT K E E L	660
TGTCATATTT S Y L	ACCCGTTATT P V I	AAACTTTTTA K L F K		TGTATACAAC V Y N	GGCTTGATTG G L I G	720
GCCTATTCCT L F L	CCTTTATGGG L Y G	TTATATATTT L Y I S	CACAGAATCA Q N Q	AGAAATTGTA E I V	GCTGTTTTTT A V F L	780
TAATCAATGT I N V	ATTGCTAGTT L L V	GCTATTTATG A I Y G		AGTTGATAAA V D K	AAAATCTTAT K I L L	840
TAAAACAGGG K Q G	TGGTTTACCT G L P	ATATTAGCTC I L A L	TTTTAACATT L T F	CTTATTTTAA L F	TACTACTTAG	900
CCGTTCGATT	TAGTTGAACG	GCTTTTAGTA	ATCATTTTT	TCTCATAATA	CAGGTAGTTT	960
AAGTAATTTG	TCTTTAAAAA	TAGTATAATA	TAACTACGAA	TTCAAAGAGA	GGTGACTTTG	1020
MTE		ACATACTAAA H T K			TCGTGTCGTT R V V	1080
GGTCAAGGTC G Q G Q		TTTTTTACAT F L H		TAAGTAGTCG S S R	CTATTTTGAT Y F D	1140
		TAAGTATTAC K Y Y			TAGAGGGCAT R G H	1200
GGCAAAAGTC G K S H		AAATACCATT N T I			TGACTTAAAG D L K	1260

GATATCTTAG TTCATTTAGA GATTGATAAA GTTATATTGG TAG	GCCATAG CGATGGTGCC 1320 G H S D G A
AATTTAGCTT TAGTTTTTCA AACGATGTTT CCAGGTATGG TTA N L A L V F Q T M F P G M V A	AGAGGGCT TTTGCTTAAT 1380 R G L L N
TCAGGGAACC TGACTATTCA TGGTCAGCGA TGGTGGGATA TTC S G N L T I H G Q R W W D I I	2110
TATAAATTCC TTCACTATTT AGGGAAACTC TTTCCGTATA TGAY K F L H Y L G K L F P Y M E	AGGCAAAA AGCTCAAGTT 1500 R Q K A Q V
ATTTCGCTTA TGTTGGAGGA TTTGAAGATT AGTCCAGCTG ATT	TTACAGCA TGTGTCAACT 1560 L Q H V S T
CCTGTAATGG TTTTGGTTGG AAATAAGGAC ATAATTAAGT TAA	AATCATTC TAAGAAACTT 1620 N H S K K L
ASYFPRGEFYSLVGI	TTTGGGCA TCACATTATT 1680 F G H H I I
AAGCAAGATT CCCATGTTTT TAATATTATT GCAAAAAAGT TTA K Q D S H V F N I I A K K F I	NDTLK
GGAGAAATTG TTGAAAAAGC TAATTGAAAA AGTCAAATCA CTO G E I V E K A N	
TGTATTTTT ATATCTGTTT TAGTGCTTAT TATTGTTGAA ATO	FATTCATT TGAAACGAAC 1860 I H L K R T
1	>
ISVEQLKSVFGQLS	CCAATGA ATCTTTTCTT 1920 PMN LF L
I S V E Q L K S V F G Q L S AATTATCCTT GTGGGGGTTA TCGCTGTCTT ACCGACAACC GGZ I I L V G V I A V L P T T G	P M N L F L ATATGACT TTGTACTGAA 1980 Y D F V L N
I S V E Q L K S V F G Q L S AATTATCCTT GTGGGGGTTA TCGCTGTCTT ACCGACAACC GGA I I L V G V I A V L P T T G TGGACTTTTA CGTACAGATA AAAGCAAAAG GTATATTTTA CAG G L L R T D K S K R Y I L Q	P M N L F L ATATGACT TTGTACTGAA 1980 Y D F V L N GACTAGTT GGTGTATCAA 2040 T S W C I N
I S V E Q L K S V F G Q L S AATTATCCTT GTGGGGGTTA TCGCTGTCTT ACCGACAACC GGA I I L V G V I A V L P T T G TGGACTTTTA CGTACAGATA AAAGCAAAAG GTATATTTTA CAG G L L R T D K S K R Y I L Q CACTTTTAAT AACTTGTCAG GATTCGGTGG CTTAATCGAT ATT T F N N L S G F G G L I D I	P M N L F L ATATGACT TTGTACTGAA 1980 Y D F V L N GACTAGTT GGTGTATCAA 2040 T S W C I N GGGGTTGC GCATGGCTTT 2100 G L R M A F
I S V E Q L K S V F G Q L S AATTATCCTT GTGGGGGTTA TCGCTGTCTT ACCGACAACC GGA I I L V G V I A V L P T T G TGGACTTTTA CGTACAGATA AAAGCAAAAG GTATATTTTA CAC G L L R T D K S K R Y I L Q CACTTTTAAT AACTTGTCAG GATTCGGTGG CTTAATCGAT ATT T F N N L S G F G G L I D I TTATGGTAAA AAAGGTCAAG AGAAGAGTGA CCTAAGAGAA GTC Y G K K G Q E K S D L R E V	P M N L F L ATATGACT TTGTACTGAA 1980 Y D F V L N GACTAGTT GGTGTATCAA 2040 T S W C I N GGGGTTGC GCATGGCTTT 2100 G L R M A F GACTCGTT TTTTACCCTA 2160 T R F L P Y
I S V E Q L K S V F G Q L S AATTATCCTT GTGGGGGTTA TCGCTGTCTT ACCGACAACC GGA I I L V G V I A V L P T T G TGGACTTTTA CGTACAGATA AAAGCAAAAG GTATATTTTA CAC G L L R T D K S K R Y I L Q CACTTTTAAT AACTTGTCAG GATTCGGTGG CTTAATCGAT ATT T F N N L S G F G G L I D I TTATGGTAAA AAAGGTCAAG AGAAGAGTGA CCTAAGAGAA GTC Y G K K G Q E K S D L R E V TCTTATTTCT GGTCTGTCAT TTATTAGTGT GATTGCCTTA ATC L I S G L S F I S V I A L I	P M N L F L ATATGACT TTGTACTGAA 1980 Y D F V L N GACTAGTT GGTGTATCAA 2040 T S W C I N GGGTTGC GCATGGCTTT 2100 G L R M A F GACTCGTT TTTTACCCTA 2160 T R F L P Y CATGAGCC ATATTTTCA 2220 M S H I F H
AATTATCCTT GTGGGGGTTA TCGCTGTCTT ACCGACAACC GGAT I I L V G V I A V L P T T G TGGACTTTA CGTACAGATA AAAGCAAAAG GTATATTTA CAGG L L R T D K S K R Y I L Q CACTTTTAAT AACTTGTCAG GATTCGGTGG CTTAATCGAT ATT T F N N L S G F G G L I D I TTATGGTAAA AAAGGTCAAG AGAAGAGTGA CCTAAGAGAA GTG Y G K K G Q E K S D L R E V TCTTATTTCT GGTCTGTCAT TTATTAGTGT GATTGCCTTA ATG L I S G L S F I S V I A L I TGCCAAAGCT AGTGTTGATT ACTATTATTT GGTATTAATT GGTATTAATTA	P M N L F L ATATGACT TTGTACTGAA 1980 Y D F V L N GACTAGTT GGTGTATCAA 2040 T S W C I N GGGGTTGC GCATGGCTTT 2100 G L R M A F GACTCGTT TTTTACCCTA 2160 T R F L P Y CATGAGCC ATATTTTCA 2220 M S H I F H GGCTAGTA TGTATTTCC 2280 A S M Y F P
AATTATCCTT GTGGGGGTTA TCGCTGTCTT ACCGACAACC GGAT I I L V G V I A V L P T T G TGGACTTTTA CGTACAGATA AAAGCAAAAG GTATATTTTA CAGG L L R T D K S K R Y I L Q CACTTTTAAT AACTTGTCAG GATTCGGTGG CTTAATCGAT ATT T F N N L S G F G G L I D I TTATGGTAAA AAAGGTCAAG AGAAGAGTGA CCTAAGAGAA GTG Y G K K G Q E K S D L R E V TCTTATTTCT GGTCTGTCAT TTATTAGTGT GATTGCCTTA ATG L I S G L S F I S V I A L I TGCCAAAGCT AGTGTTGATT ACTATTATTT GGTATTAATT GGTATTATTT TTGCCAAAGCT AGTGTTGATT ACTATTATTT TTGCTATTTATTATT TTGCTATTTATTATT TTGCTATTTTATTT	PMNLFL ATATGACT TTGTACTGAA 1980 YDFVLN GACTAGTT GGTGTATCAA 2040 TSWCIN GGGTTGC GCATGGCTTT 2100 GLRMAF GACTCGTT TTTTACCCTA 2160 TRFLPY CATGAGCC ATATTTTCA 2220 MSHIFH GGCTAGTA TGTATTTTCC 2280 ASMYFP CGGGAGATA TGCCATCTAG 2340 GDMPSS
AATTATCCTT GTGGGGGTTA TCGCTGTCTT ACCGACAACC GGAT I I L V G V I A V L P T T G TGGACTTTTA CGTACAGATA AAAGCAAAAG GTATATTTTA CAGG L L R T D K S K R Y I L Q CACTTTTAAT AACTTGTCAG GATTCGGTGG CTTAATCGAT ATT T F N N L S G F G G L I D I TTATGGTAAA AAAGGTCAAG AGAAGAGTGA CCTAAGAGAA GTOY G K K G Q E K S D L R E V TCTTATTTCT GGTCTGTCAT TTATTAGTGT GATTGCCTTA ATG L I S G L S F I S V I A L I TGCCAAAGCT AGTGTTGATT ACTATTATTT GGTATTAATT GGTATTATTTATTATTT TGGATTTCT GTCATTATTT TTGTTATTTATTT TTGTTATTTATTT TTGTTATTTAT	P M N L F L ATATGACT TTGTACTGAA 1980 Y D F V L N GACTAGTT GGTGTATCAA 2040 T S W C I N GGGGTTGC GCATGGCTTT 2100 G L R M A F GACTCGTT TTTTACCCTA 2160 T R F L P Y CATGAGCC ATATTTTTCA 2220 M S H I F H GGCTAGTA TGTATTTTCC 2280 A S M Y F P CGGAGATA TGCCATCTAG 2340 G D M P S S ATGTGCGG CCGCAGCATT 2400 C A A A A F

0 1	G C	A	V G	I	TGTATC V S	CCTT!	ATTCC(I P	C GG1 G	'GGAT' G L	TAG G	GAAGT S	rttt F	'GA E	2520
ATTAGT L V	TCTA L F	TTTAC. T	AGGGT G F	TTGC' A	TGCCGA A E	GGGA	CTACC! L P	r aa <i>i</i> K	AGAAA E T	CTG V	TGGTT V		NTG W	2580
GTTATT L L	ractt y	TATCG R	TTTAG L A		CTATAT Y I		CCATTO P F	C TTT	rgcag A g	GTA I	TCTAT Y	TTTC F	TT F	2640
TATCCA I H	Y L	TTAGG G	TAGTC S Q	AAAT. I	AAATCA N Q		TATGA Y E	A AAT N	rgtcc V p	CGA K	AAGA(E		GT V	2700
ATCAAC S T	V L	CTACA Q	AACCA T M	TGGT V	GAGCCA S H		ATGCG M R	r ATT I >	TTAG L G	GTG A	CATT(AT I	2760
ATTTTC F S	CAACA T A		TTTTG F E	AAAA' N	TATTAC I T		ATTATO	G TGO W	STTGC: L Q	AGA K	AGCTA L	AGGC G	TT L	2820
GGACCO D P	CATTA (CAAGA E	ACAAA Q M	TGTT.	ATGGCA W Q		CCAGG' P G	r TT <i>I</i> L	ATTGC' L L	IGG G	GGGT7	TTGT C	TT F	2880
TATTCT I L	L A	R	AACTA T I	D	TCAAAA Q K	V	K N	A	TTTTC F P	I	TTGCT A	TTAT I	I	2940
CTGGAT W I	TACT T L				TCTTAA L N							ATCT S		3000
W F	I L	L	L L	G	CTTATT L L	V :	I K	P	T L	Y	K	K	Q	3060
ATTTAT F I			GGAAG E E	AGCG R	TATTAA I K			C ATT	TATCG' I V	rta S	GTTTA L		GG G	3120
V L			TGCAG A G		ACTATT L F		ATCAGO I R	G GCI A	CATA' H I		CAGGT G		'AG S	3180
V L TATTGA	F Y AACGC R L	I CTGCA H	A G TTATA Y I	L TCAT. I	L F AGCATG A W	P GGAG E	I R CCGATA P I	A A GCA A	H I ATTGG L A	T CTA T	G CGTT(L	G SATT I	S CT L	3180 3240
V L TATTGA I E TACTCA T L	F Y AACGC R L CGTT V Y	I CTGCA H TATTT. L	A G TTATA Y I ATGTT C L	TCAT. I TGGT	L F AGCATG A W TAAGAT K I	GGAGGE TTTAG	I R CCGATA P I CAAGGA Q G	A GCA A A A AAA	H I ATTGG L A ATCTT	T CTA T GTC Q	G CGTTC L AGATT	G EATT I IGGT G	S CT L 'GA D	
V L TATTGA I E TACTCT T L TGTGTT V F	F Y AACGC R L ICGTT V Y ICAAT N V	I CTGCA H TATTT L GTGGA D	A G TTATA Y I ATGTT C L TCGTT R Y	TCAT. I TGGT V ATAA. K	L F AGCATG A W TAAGAT K I AAAACT K L	GGAGGE TTTAGL ACTTGL	I R CCGATA P I CAAGGA Q G CAAGCA	A GCA A AAA K T TAC	H I ATTGG L A ATCTT S C CGGTG G G	T CTA T GTC Q GTT S	G CGTTC L AGATT I CTTCC	G EATT I IGGT G GAT	S CT L GA D 'AG S	3240
V L TATTGA I E TACTCA T L TGTGTA V F CGGTTA G L	F Y AACGC R L ICGTT V Y ICAAT N V IAGCC A F	I CTGCA H TATTT L GTGGA D TTTTT	A G TTATA Y I ATGTT C L TCGTT R Y AAATG N D	TCAT. I TGGT V ATAA. K ATAA.	L F AGCATG A W TAAGAT K I AAAACT K L AAGGCT R L	GGAGGE TTTAG L ACTTG L CTAC'	I R CCGATA P I CAAGGA Q G CAAGCA Q A TGGTAGW Y	A GCA A AAA K T TAC Y C CAA	H I ATTGG L A ATCTT S C CGGTG G G AAAAA K N	T CTA T GTC Q GTT S ATG	G CGTTC L AGATT I CTTCC S GAGAA	G EATT I EGGT D EGAT D	S CT L GA D AG S TG C	32 4 0 3300
V L TATTGA I E TACTCT T L TGTGTT V F CGGTTT G L CGTTGC V A	F Y AACGC R L ICGTT V Y ICAAT N V IAGCC A F CGTTC F Q	I CTGCA H TATTT L GTGGA D TTTTT L CAATT F	TTATA Y I ATGTT C TCGTT R Y AAATG N D TGTAA V I	TCAT. I TGGT V ATAA K ATAA K TTGT V	L F AGCATG A W TAAGAT K I AAAACT K L AAGGCT R L CAATAA N N	GGAGGE TTTAGL ACTTGL CTACTY TAAAT	I R CCGATA P I CAAGGA Q G CAAGCA Q A TGGTACA W Y TGTCTA	A GCAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA	H I ATTGG L A ATCTT S C CGGTG G G AAAAA K N TATGG	T CTA T GTC GTT S ATG GGG	GCTTCC LAGATT ICTTCC SCAGAN EAACCN P	G EATT I IGGT G EGAT D AGAT AGCC	S CT L GA D AG S TG C	3240 3300 3360 3420
TATTGATE TACTCT T L TGTGTT V F CGGTTT G L CGTTGC V A TGATGAT D D	F Y AACGC R L ICGTT V Y ICAAT N V IAGCC A F CGTTC F Q ACACT T Y	I CTGCA H TATTT L GTGGA D TTTTT L CAATT F TATAT I	TTATA Y I ATGTT C I TCGTT R Y AAATG N D TGTAA V I TCGTG R E	TCAT. I TGGT V ATAA K ATAA K TTGT V AAGC A	AGCATGA W TAAGATKI AAAACTKL AAGGCTRL CAATAANN TATTGA	GGAGGE TTTAGL ACTTGL CTAC' Y TAAA' K ATCG'	I R CCGATA P I CAAGGA Q G CAAGC' Q A TGGTAC W Y TGTCT' C L TTTAT' F I	A GCAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA	H I ATTGG L A ATCTT S C CGGTG G G AAAAA K N TATGG M G TGATG	T CTA GTC GTTS ATG GGG E CTG	GCGTTCC LAGATT ICTTCC SCAGATE AACCT PATAACK	G EATT I EGGT D AGAT AGCO AGCO A	S CT L GA D AG S C GG G GA	3240 3300 3360 3420 3540
TATTGATE TACTCT T L TGTGTT V F CGGTTT G L CGTTGC V A TGATGAT D D	F Y AACGC R L CGTT V Y CAAT N V CAAT N V CGTTC F Q ACACT T Y ACCTT L V	CTGCA H TATTT L GTGGA D TTTTT L CAATT F TATAT I GTTTT F	TTATA Y I ATGTT C L TCGTT R Y AAATG N D TGTAA V I TCGTG R E TTACA Y S	TCAT. I TGGT V ATAA. K ATAA. K TTGT V AAGC. A GTAT. I	L F AGCATG A W TAAGAT K I AAAACT K L AAGGCT R L CAATAA N N TATTGA I E TGGACA G Q	GGAGGE TTTAGL ACTTGL CTAC' Y TAAA' K ATCG' S GAAG' K	I R CCGATA P I CAAGGA Q G CAAGCA Q A TGGTACA W Y TGTCTA TTTAT F I TTGACA L T	A GCAA A AAA K TAC Y C CAA Q T ATT I GAT A CTA	H I ATTGG L A ATCTT S C CGGTG G G AAAAA K N CATGG M G CGATG D A ACTTT L L	TTA GTQ GTTS ATG GGG TAC TAC H	GCGTTC L AGATT I CTTCC S GAGAME AACCM P ATAACM K ATGACME	G SATT I GGT GGAT D AGCO AGCO A CTA L STAT	S CT L GA D AG S C GG G G G G G	3240 3300 3360 3420

G N K Y K P F	R N A	L N R V E K D G F Y	3720
TTTCGAAGTT GTACAATCGC F E V V Q S P	CACATAGTCA H S Q	AGAGCTACTA AATAGTTTGG AAGAGATTTC E L L N S L E E I S	3780
TAATACTTGG TTAGAAGGAC N T W L E G R		AGGTTTCTCA CTAGGATATT TTAATAAAGA G F S L G Y F N K D	3840
TTATTTCCAA CAAGCCCCAA Y F Q Q A P I	TAGCTTTGGT A L V	AAAAAATGCT GAACACGAAG TTGTTGCTTT K N A E H E V V A F	3900
TGCTAATATT ATGCCAAACT A N I M P N Y	ATGAAAAGAG E K S	TATTATCTCT ATTGATTTAA TGCGTCACGA I I S I D L M R H D	3960
TAAACAGAAA ATTCCGAATG K Q K I P N G	GCGTTATGGA V M D	TTTCCTCTTT TTATCATTAT TCTCTTATTA F L F L S L F S Y Y	4020
TCAAGAGAAG GGATACCACT Q E K G Y H Y	ATTTTGATTT F D L	GGGGATGGCA CCTTTATCAG GAGTTGGTCG G M A P L S G V G R	4080
CGTTGAAACA AGTTTTGCTA V E T S F A K	AAGAGAGAAT E R M	GGCGTATCTT GTCTATCATT TCGGTAGTCA A Y L V Y H F G S H	4140
TTTCTACTCA TTTAATGGTT F Y S F N G L	TACACAAGTA H K Y	TAAGAAGAAG TTTACACCAT TGTGGTCGGA K K K F T P L W S E	4200
ACGTTATATT TCTTGTTCTC R Y I S C S R		GTTAATTTGT GCTATTTGTG CCCTATTAAT L I C A I C A L L M	4260
GGAAGATAGT AAAATTAAGA E D S K I K I		AGCTTTATTT GGCAATTAAA AAGAGCATGT	4320
CATGCGACAT GCTCTTTTA	AATCATTTAA	TACCATTGAT TGCTTGAATC TACTTTATAA	4380
TATGATGTGC TTTTAAATAT	TGTTTAGCTA	CTGTAGCTGC TGATTTATGC TTTACAGCTA	4440
CTTGGTAGTT CATTTCTTGC	ATTTCTTTTT	CAGTGATATG ACCAGCAAGT TTATTGAGAG	4500
CTTTTTTTAC TTGA (SEQ	ID NO:1)		4514

FIG. 1a [clonel-dna/aa]

SGKEPANRFS	WAKNKLLING	FIATLAATIL	FFAVQFIGLK	PDYPGKTYFI	50
ILLTAWTLMA	LVTALVGWDN	RYGSFLSLLI	LLFQLGSSAG	TYPIELSPKF	100
FQTIQPFLPM	TYSVSGLRET	ISLTGDVNHQ	WRMLVIFLVS	SMILALLIYR	150
KQED (SEQ	ID NO:2)				154
		FIG. 1	b		
MSTLTIIIAT	LTALEHFYIM	YLETLATQSN	MTGKIFSMSK	EELSYLPVIK	50
LFKNQGVYNG	LIGLFLLYGL	YISQNQEIVA	VFLINVLLVA	IYGALTVDKK	100
ILLKQGGLPI	LALLTFLF (SEQ ID NO:3	3)		118
		FIG. 1	C		
MTENWLHTKD	GSDIYYRVVG (OGOPTVFT.HG	NST.SSRVEDK	OTAVECEVVO	5 0
					50
	KSHAKLNTIS I				100
LALVFQTMFP	GMVRGLLLNS (GNLTIHGQRW	WDILLVRIAY	KFLHYLGKLF	150
PYMRQKAQVI	SLMLEDLKIS I	PADLQHVSTP	VMVLVGNKDI	IKLNHSKKLA	200

FIG. 1d

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SYFPRGEFYS LVGFGHHIIK QDSHVFNIIA KKFINDTLKG EIVEKAN

(SEQ ID NO:4)

MIHLKRTISV	EQLKSVFGQL	SPMNLFLIIL	VGVIAVLPTT	GYDFVLNGLL	50
RTDKSKRYIL	QTSWCINTFN	NLSGFGGLID	IGLRMAFYGK	KGQEKSDLRE	100
VTRFLPYLIS	GLSFISVIAL	IMSHIFHAKA	SVDYYYLVLI	GASMYFPVIY	150
WISGHKGSHY	FGDMPSSTRI	KLGVVSFFEW	GCAAAAFIII	GYLMGIHLPV	200
YKILPLFCIG	CAVGIVSLIP	GGLGSFELVL	FTGFAAEGLP	KETVVAWLLL	250
YRLAYYIIPF	FAGIYFFIHY	LGSQINQRYE	NVPKELVSTV	LQTMVSHLMR	300
ILGAFLIFST	AFFENITYIM	WLQKLGLDPL	QEQMLWQFPG	LLLGVCFILL	350
ARTIDQKVKN	AFPIAIIWIT	LTLFYLNLGH	ISWRLSFWFI	LLLLGLLVIK	400
PTLYKKQFIY	SWEERIKDGI	IIVSLMGVLF	YIAGLLFPIR	AHITGGSIER	450
LHYIIAWEPI	ALATLILTLV	YLCLVKILQG	KSCQIGDVFN	VDRYKKLLQA	500
YGGSSDSGLA	FLNDKRLYWY	QKNGEDCVAF	QFVIVNNKCL	IMGEPAGDDT	550
YIREAIESFI	DDADKLDYDL	VFYSIGQKLT	LLLHEYGFDF	MKVGEDALVN	600
LETFTLKGNK	YKPFRNALNR	VEKDGFYFEV	VQSPHSQELL	NSLEEISNTW	650
LEGRPEKGFS	LGYFNKDYFQ	QAPIALVKNA	EHEVVAFANI	MPNYEKSIIS	700
IDLMRHDKQK	IPNGVMDFLF	LSLFSYYQEK	GYHYFDLGMA	PLSGVGRVET	750
SFAKERMAYL	VYHFGSHFYS	FNGLHKYKKK	FTPLWSERYI	SCSRSSWLIC	800
AICALLMEDS	KIKIVK (SE	EQ ID NO:5)			816

FIG. 1e

MRILGAFLIF	STAFFENITY	IMWLQKLGLD	PLQEQMLWQF	PGLLLGVCFI	50
LLARTIDQKV	KNAFPIAIIW	ITLTLFYLNL	GHISWRLSFW	FILLLGLLV	100
IKPTLYKKQF	IYSWEERIKD	GIIIVSLMGV	LFYIAGLLFP	IRAHITGGSI	150
ERLHYIIAWE	PIALATLILT	LVYLCLVKIL	QGKSCQIGDV	FNVDRYKKLL	200
QAYGGSSDSG	LAFLNDKRLY	WYQKNGEDCV	AFQFVIVNNK	CLIMGEPAGD	250
DTYIREAIES	FIDDADKLDY	DLVFYSIGQK	LTLLHEYGF	DFMKVGEDAL	300
VNLETFTLKG	NKYKPFRNAL	NRVEKDGFYF	EVVQSPHSQE	LLNSLEEISN	350
TWLEGRPEKG	FSLGYFNKDY	FQQAPIALVK	NAEHEVVAFA	NIMPNYEKSI	400
ISIDLMRHDK	QKIPNGVMDF	LFLSLFSYYQ	EKGYHYFDLG	MAPLSGVGRV	450
ETSFAKERMA	YLVYHFGSHF	YSFNGLHKYK	KKFTPLWSER	YISCSRSSWL	500
ICAICALLME	DSKIKIVK	(SEQ ID NO	: 6)		518

FIG. 1f

AATTTTGATA TCGAAACAAC N F D I E T T	AACTTTTGAG GCAA T F E A M	TGAAAA AGCACGCGTC I K K H A S	ATTATTGGAG L L E	60
AAAATATCTG TTGAGCGTTC K I S V E R S	TTTTATTGAA TTTG F I E F D	ATAAAC TTCTATTAGO K L L L A	ACCTTATTGG P Y W	120
CGTAAAGGAA TGCTGGCACT R K G M L A L	AATAGATAGT CATG		ATGCTTAAAA C L K	180
AATAGGGAAT TACAATTAAG N R E L Q L S	CGCCTTTTTG TCCC		TTTATTTGAG L F E	240
ACATCAGAAC AAGCTTGGGC T S E Q A W A	ATCACTCATC TTGA S L I L S		CACAAAGACT T K T	300
TTTTTAAAAA AATGGAAGAC F L K K W K T	ATCAACTCAC TTTC S T H F Q		TATAGTGGAT I V D	360
GTTTATCGTA TTCGTGAACA V Y R I R E Q	AATGGGATTG GCTA M G L A K		TTATGGAAAA Y G K	420
ACTATAATAA AACAAGCGGA T I I K Q A E	AGGTATTCGC AAAG G I R K A	CAAGAG GCTTGATGGT R G L M V	TGATTTCGAA D F E	480
AAAATAGAAC AACTAGATAG K I E Q L D S	TGAGTTAGCA ATCC E L A I H		AGTTGTCAAT V V N	540
GGTGGCACCT TAATCAAGAA G G T L I K K	ATTAGGAATA AAAC L G I K P		AGATATTATC D I I	600
TCTCAAATTG AATTAGCCAT S Q I E L A I	TGTTTTAGGA CAAC V L G Q L	TGATTA ATGAAGAAGA I N E E E	GGCTATTTTA A I L	660
CATTTTGTTA AGCAGTACTT H F V K Q Y L	GATGGATTAG AGAG M D	GATTAT ATGAGCGATT	TTTTAGTAGA L V D	720
TGGATTGACT AAGTCGGTTG G L T K S V G		TTTAGT AATGTTTCAT F S N V S F	TTATCATCCA I I H	780
TAGTTTAGAC CGTATTGGGA S L D R I G I		GGAACT GGAAAGACAA G T G K T T	CACTATTAGA L L D	840
TGTTATTTCG GGTGAATTAG V I S G E L G		CGTTCC CCTTTTTCAT R S P F S S	CAGCTAATGA A N D	900
TTATAAGATT GCTTATTTAA Y K I A Y L K		TTTGAT GATTCTCAGA F D D S Q T	CAATTTTGGA I L D	960
CACCGTACTT TCTTCTGACT	TAAGAGAGAT GGCT R E M A		AATTATTGCT L L L	1020
TAATCACTAC GAAGAAAGTA N H Y E E S K	AGCAATCACG TCTA Q S R L		AAATGGATTC M D S	1080
TTTAGATGCT TGGTCTATTG L D A W S I E				1140
TGATTTGCAG TTGTCGGTTG D L Q L S V G	GTGAATTATC AGGA E L S G		TTCAATTAGC Q L A	1200

	TTAAATGATG L N D A	CAGATTTATT D L L	GCTCTTAGAC L L D	GAACCTACTA E P T N	ACCACTTAGA H L D	1260
	ATTGCATGGT I A W L	TAACGAATTT T N F	TTTGAAAAAT L K N	AGTAAAAAGA S K K T	CAGTGCTTTT V L F	1320
	GATCGTTATT O R Y F	TTCTAGACAA L D N	TGTTGCAACA V A T	CGTATTTTTG R I F E	AATTAGATAA L D K	1380
AQIT	T E Y Q	AAGGCAATTA G N Y	Q D Y	V R L R	A E Q	1440
DERD	O A A S	GTTTACATAA L H K	K K Q	L Y K Q	E L A	1500
TTGGATGCGT A W M R I	r Q P Q	ARA	T K Q	Q A R I	N R F	1560
TCAAAATCTA A Q N L K	KNDL	H Q T	S D T	S D L E	M T F	1620
E T S R	RIGK	AAAAGGTTAT K V I			TTTCTTACCC S Y P	1680
D K S I	I L K D	F N L	L I Q	AATAAAGACC N K D R	GTATTGGCAT I G I	1740
VGDN	1 G V G	GAAAGTCAAC K S T	L L N	TTAATTGTTC L I V Q	AAGATTTACA D L Q	1800
P D S G	S N V S	I G E	TIR	GTAGGTTACT V G Y F	TTTCACAACA S Q Q	1860
L H N M	1 D G S	K R V	I N Y	L Q E V	TTGCAGATGA A D E	1920
V K T S	S V G T	CAACAAGTGT T S V	T E L	TTGGAACAAT L E Q F	TTCTCTTTCC L F P	1980
	H G T Q	I A K	L S G	GGTGAGAAA G E K K	AAAGACTTTA R L Y	2040
r r k i	LIE	AAAAGCCTAA K P N	V L L	CTTGATGAGC L D E P	CGACAAATGA T N D	2100
CTTAGATATT G L D I A	A T L T	V L E	N F L	Q G F G	G P V	2160
	S H D R	Y F L	D K V	A N K I	I A F	2220
	DIRE	F F G	N Y T	D Y L D	E K A	2280
~	NNE	V I S	K K E	S T K T	S R E	2340
_	RKRM	S Y F	E K Q	E W A T	I E D	2400
CGATATTATG A	TALIGGAAA	ATACTATCAC	TCGTATAGAA	AATGATATGC	AAACATGTGG	2460

D	I	M	I	L	E	N	T	I	T	R	I	E		N D	M	Q I	T	С	G	
TAC	STG# D	ATTTT F	ACZ T		GTI L	TAT S	CTG		TACA Q	AAZ K			'A	GATG D A		_	ATG.		CACT L	2520
TCT L		AAAAG K	TA: Y	rga D	CCC R	TT Y	ATG/ E	AGTI Y	ACCT L	TAG	GTG/ E	AGTT. L	'A	GACA D T		GAT I	TAT	CCG' R	rccg P	2580
ATT I	rtan I	TAAAA K N			GAC D	CCA Q	AGCI A		rgca A		ATTA L		'C R	GACA Q	AAG S	TTT L	ACG R	CGC(CTAT Y	2640
GAT D	ltta L	AGATA D K			GAI D	TAC T	AGCI A	ATA: Y	rtca s	GA(CCC' P		T	TAGA D	TCA H	TTT L	GAC T	CTC S	ATAC Y	2700
TAC Y	CGAA E	K I			AAC K	STC S	AGGI G	ATT(F	CTTT F	GT(V	CAT'		G E	AGAG. R	AGA D	TGA E	GAT'	TAT'	rggc G	2760
TG1 C	rggc G	CGGCT G F			CCG P	CT L	GAAI K	AAA: N	ICTA L	AT:	rgcz A		A M	TGCĄ	GAA K	GGT V	GTA Y	CAT'	rgca A	2820
GA/ E	ACGI R	TTTCC F R			AAC K	G G	GCT:	rgc: A	TACT T	GA:	rtti L		A K	AAAT M	GAT I	TGA E	AGT. V	AGAZ E	AGCT A	2880
CGI R	AAA <i>I</i> K	ATTG I G			AG <i>I</i> R	ACA Q	ACT:	TTA: Y	L L	GA0 E	GAC T		A S	GTAC' T	TTT L	GAG S	TAG R	GGCI A	AACT T	2940
GC0 A	GTI V	TATA Y K			ATC M	G G	ATA'	rtg: C	IGCC A	TTA L	ATC S		C P	CAAT	AGC A	AAA N	TGA' D	TCAI Q	AGGT G	3000
CAT H	rac <i>i</i> T	AGCTA A M		GAT D	'AT'I	TTG W	GAT(GAT:	raaa K		rtt: L	ATAA	.G	TTGA	AAG	TGG	ATT	AGT(GAAC	3060
ATO	GGAI	TAAT	TA	ГТТ	'TG <i>I</i>	AGA	TAA	GAG	GAAA	GA	AAA	GGAG.	A	CATA	TAT M _	GGC A	ATA Y	TAT'	rtgg W	3120
TC: S	rati Y	TTGA L K				CCC P	CAA'	TTG(W	GTTA L	TG(W	GCT' L	TGAT D	T L	TACT.	AGG G	AGC A	TAT	GCT: L	rttt F	3180
GT(V	GACO T	GTTA V I			GGI G	TA <i>I</i> M	GCC(CACI T	AGCC A	TTI L	AGC(A		'A M	TGAT	TGA D	TAA N	TGG G	CGT: V	TACA T	3240
AA! K	AGG1 G	TGATC D R			'GG <i>I</i> G	AGT V	TTA;		GTGG W	ACC T	GTT(F		A M	TGTT F		ATT F	TGT V		ACTA L	3300
GG:	rati I	TATTG I G		CGT R	'ATT	TAC T	GAT(GGC'. A	TTAC Y	GCI A	ATC' S		C R	GCTT.	AAC T	GAC T	AAC. T	AAT(M	GATT I	3360
	AGA1 D	TATGC M R	GT2	AAT N		TAT M	GTA'				rca. Q		T Y	ACTC S		TCA H		ATA' .Y		3420
	GAT <i>I</i> I	AGGTG G V			TC# S		AGT V		ACGT R		GAC T		G D		TTT F	TGT V		GAT M	GCAA Q	3480
		rgaaa E M			TT# L		TTT:		CCTA L	GTZ V	AAC' T		'A M	TGGT. V	AAT M	GAT I	TTT	TAG S	CGTG V	3540
GT: V		SATAC I L				SAG S	TCC:		TTTG L	GC' A			G V	TAGC A	GGT V	TGC A		GCC' P		3600
		AGGAG G V							TATA					CTTT		TGA		ACA		3660

ACTATGCTTG ATAAAATCAA	TCAATATGTT CGTGAAAATT	TAACAGGGTT ACGCGTTGTT	3720
T M L D K I N	Q Y V R E N L	T G L R V V	
AGAGCCTTTG CAAGAGAGAA	TTTTCAATCA CAAAAATTTC	AAGTCGCTAA CCAACGTTAC	3780
R A F A R E N	F Q S Q K F Q	V A N Q R Y	
ACAGATACTT CAACTGGTCT	TTTTAAATTA ACAGGGCTAA	CAGAACCACT TTTCGTTCAA	3840
T D T S T G L	F K L T G L T	E P L F V Q	
ATTATTATTG CAATGATTGT I I I A M I V	GGCTATCGTT TGGTTTGCTT A I V W F A L	TGGATCCCTT ACAAAGAGGT D P L Q R G	3900
GCTATTAAAA TAGGGGATTT	AGTTGCTTTT ATCGAATATA	GCTTCCATGC TCTCTTTTCA	3960
A I K I G D L	V A F I E Y S	F H A L F S	
TTTTTGCTAT TTGCCAATCT	TTTTACTATG TATCCTCGTA	TGGTGGTATC AAGCCATCGT	4020
F L L F A N L	F T M Y P R M	V V S S H R	
ATTAGAGAGG TGATGGATAT	GCCAATCTCT ATCAATCCTA	ATGCCGAAGG TGTTACGGAT	4080
I R E V M D M	P I S I N P N	A E G V T D	
ACGAAACTTA AAGGGCATTT	AGAATTTGAT AATGTAACAT	TCGCTTATCC AGGAGAAACA	4140
T K L K G H L	E F D N V T F	A Y P G E T	
GAGAGTCCCG TTTTGCATGA	TATTTCTTTT AAAGCTAAGC	CTGGAGAAAC AATTGCTTTT	4200
E S P V L H D	I S F K A K P	G E T I A F	
ATTGGTTCAA CAGGTTCAGG I G S T G S G	AAAATCTTCT CTTGTTAATT K S S L V N L	TGATTCCACG TTTTTATGAT I P R F Y D	4260
GȚGACACTTG GAAAAATCTT	AGTAGATGGA GTTGATGTAA	GAGATTATAA CCTTAAATCA	4320
V T L G K I L	V D G V D V R	D Y N L K S	
CTTCGCCAAA AGATTGGATT L R Q K I G F	TATCCCCCAA AAAGCTCTTT I P Q K A L L		4380
GAGAATTTAA AATATGGAAA	AGCTGATGCT ACTATTGATG	ATCTTAGACA AGCGGTTGAT	4440
E N L K Y G K	A D A T I D D	L R Q A V D	
ATTTCTCAAG CTAAAGAGTT	TATTGAGAGT CACCAAGAAG	CCTTTGAAAC GCATTTAGCT	4500
I S Q A K E F	I E S H Q E A	F E T H L A	
GAAGGTGGGA GCAATCTTTC	TGGGGGTCAA AAACAACGGT	TATCTATTGC TAGGGCTGTT	4560
E G G S N L S	G G Q K Q R L	S I A R A V	
GTTAAAGATC CAGATTTATA V K D P D L Y	TATTTTTGAT GATTCATTTT I F D D S F S	CTGCTCTCGA TTATAAGACA A L D Y K T	4620
GACGCTACTT TAAGAGCGCG D A T L R A R	TCTAAAAGAA GTAACCGGTG L K E V T G D		4680
GCTCAAAGGG TGGGTACGAT	TATGGATGCT GATCAGATTA	TTGTCCTTGA TGAAGGCGAA	4740
A Q R V G T I	M D A D Q I I	V L D E G E	
ATTGTCGGTC GTGGTACCCA	CGCTCAATTA ATAGAAAATA	ATGCTATTTA TCGTGAAATC	4800
I V G R G T H	A Q L I E N N	A I Y R E I	
GCTGAGTCAC AACTGAAGAA A E S Q L K N	CCAAAACTTA TCAGAAGGAG Q N L S E G E	AGTGATTGTA TGAGAAAAA M R K K >	4860

ATCTGTTTTT TTGAGATTAT GGTCTTACCT AACTCGCTAC AAAG	
GATTTTTTT AAAGTTTTAT CTAGTTTTAT GAGTGTTCTG GAGCO	CTTTTA TTTTAGGGTT 4980 F I L G L
AGCGATAACA GAGTTGACTG CTAACCTTGT TGATATGGCT AAGGGA A I T E L T A N L V D M A K G	
ATTGAACGTT CCTTATATTG CTGGTATTTT GATTATTTAT TTTTT	
TGAATTAGGT TCTTATGGCT CAAATT (SEQ ID NO:7)	5126

FIG. 2a

NFDIETTTFE	AMKKHASLLE	KISVERSFIE	FDKLLLAPYW	RKGMLALIDS	50
HAFNYLPCLK	NRELQLSAFL	SQLDKDFLFE	TSEQAWASLI	LSMEVEHTKT	100
FLKKWKTSTH	FQKDVEHIVD	VYRIREQMGL	AKEHLYRYGK	TIIKQAEGIR	150
KARGLMVDFE	KIEQLDSELA	IHDRHEIVVN	GGTLIKKLGI	KPGPQMGDII	200
SQIELAIVLG	QLINEEEAIL	HFVKQYLMD	(SEQ ID NO:	8)	229

FIG. 2b

Mangrander	KSVGDKTVFS	NVSFIIHSLD	RIGIIGVNGT	GKTTLLDVIS	50
GELGFDGDRS	PFSSANDYKI	AYLKQEPDFD	DSQTILDTVL	SSDLREMALI	100
KEYELLLNHY	EESKQSRLEK	VMAEMDSLDA	WSIESEVKTV	LSKLGITDLQ	150
LSVGELSGGL	RRRVQLAQVL	LNDADLLLLD	EPTNHLDIDT	IAWLTNFLKN	200
SKKTVLFITH	DRYFLDNVAT	RIFELDKAQI	TEYQGNYQDY	VRLRAEQDER	250
DAASLHKKKQ	LYKQELAWMR	TQPQARATKQ	QARINRFQNL	KNDLHQTSDT	300
SDLEMTFETS	RIGKKVINFE	NVSFSYPDKS	ILKDFNLLIQ	NKDRIGIVGD	350
NGVGKSTLLN	LIVQDLQPDS	GNVSIGETIR	VGYFSQQLHN	MDGSKRVINY	400
LQEVADEVKT	SVGTTSVTEL	LEQFLFPRST	HGTQIAKLSG	GEKKRLYLLK	450
ILIEKPNVLL	LDEPTNDLDI	ATLTVLENFL	QGFGGPVITV	SHDRYFLDKV	500
ANKIIAFEDN	DIREFFGNYT	DYLDEKAFNE	QNNEVISKKE	STKTSREKQS	550
RKRMSYFEKQ	EWATIEDDIM	ILENTITRIE	NDMQTCGSDF	TRLSDLQKEL	600
DAKNEALLEK	YDRYEYLSEL	DT (SEQ II	NO:9)		622

FIG. 2c

MIIRPIIKND DQAVAQLIRQ SLR	AYDLDKP DTAYSDPHLD HLTSYYEKIE 5	50
KSGFFVIEER DEIIGCGGFG PLK	NLIAEMQ KVYIAERFRG KGLATDLVKM 10	0 (
IEVEARKIGY RQLYLETAST LSR	ATAVYKH MGYCALSQPI ANDQGHTAMD 15	50
IWMIKDL (SEQ ID NO:10)	15	57

FIG. 2d

MAIIWSILKR	IBMMTMTDFF	GAMLEVIVIL	GMPTALAGMI	DNGVTKGDRT	50
GVYLWTFIMF	IFVVLGIIGR	ITMAYASSRL	TTTMIRDMRN	DMYAKLQEYS	100
HHEYEQIGVS	SLVTRMTSDT	FVLMQFAEMS	LRLGLVTPMV	MIFSVVMILI	150
TSPSLAWLVA	VAMPLLVGVV	LYVAIKTKPL	SERQQTMLDK	INQYVRENLT	200
GLRVVRAFAR	ENFQSQKFQV	ANQRYTDTST	GLFKLTGLTE	PLFVQIIIAM	250
IVAIVWFALD	PLQRGAIKIG	DLVAFIEYSF	HALFSFLLFA	NLFTMYPRMV	300
VSSHRIREVM	DMPISINPNA	EGVTDTKLKG	HLEFDNVTFA	YPGETESPVL	350
HDISFKAKPG	ETIAFIGSTG	SGKSSLVNLI	PRFYDVTLGK	ILVDGVDVRD	400
YNLKSLRQKI	GFIPQKALLF	TGTIGENLKY	GKADATIDDL	RQAVDISQAK	450
EFIESHQEAF	ETHLAEGGSN	LSGGQKQRLS	IARAVVKDPD	LYIFDDSFSA	500
LDYKTDATLR	ARLKEVTGDS	TVLIVAQRVG	TIMDADQIIV	LDEGEIVGRG	550
THAQLIENNA	IYREIAESQL	KNQNLSEGE	(SEQ ID NO:	11)	579

FIG. 2e

MRKKSVFLRL	WSYLTRYKAT	LFLAIFLKVL	SSFMSVLEPF	ILGLAITELT	50
ANLVDMAKGV	SGAELNVPYI	AGILIIYFFR	GVFYELGSYG	SN	92
(SEQ ID NO:	:12)				

FIG. 2f

AATTTGGAAG TGCTCTATCA	ACAGTTGAAG	TAAAGGAGAT	TATTAGTGAA	GAAAACATAT	60
F G S A L S	T V E V	K E I	I S E	E N I W	
GGTTATATCG GCTCAGTTGC	TGCCATTTTA	CTAGCTACTC	ATATTGGAAG	TTACCAACTT	120
L Y R L S C	C H F T	S Y S	Y W K	L P T W	
GGTAAGCATC ATATGGGTCT M G L	AGCAACAAAG A T K	GACAATCAGA D N Q I	TTGCCTATAT A Y I	TGATGACAGC D D S	180
AAAGGTAAGG CAAAAGCCCC	TAAAACAAAC	AAAACGATGG	ATCAAATCAG	TGCTGAAGAA	240
K G K A K A P	K T N	K T M D	Q I S	A E E	
GGCATCTCTG CTGAACAGAT	CGTAGTCAAA	ATTACTGACC	AAGGCTATGT	GACCTCACAC	300
G I S A E Q I	V V K	I T D Q	G Y V	T S H	
GGTGACCATT ATCATTTTTA	CAATGGGAAA	GTTCCTTATG	ATGCGATTAT	TAGTGAAGAG	360
G D H Y H F Y	N G K	V P Y D	A I I	S E E	
TTGTTGATGA CGGATCCTAA	TTACCGTTTT	AAACAATCAG	ACGTTATCAA	TGAAATCTTA	420
L L M T D P N	Y R F	K Q S D	V I N	E I L	
_	CAATGGCAAC N G N	TATTATGTTT Y Y V Y		AGGTAGTAAG G S K	480
CGCAAAAACA TTCGAACCAA	ACAACAAATT	GCTGAGCAAG	TAGCCAAAGG	AACTAAAGAA	540
R K N I R T K	Q Q I	A E Q V	A K G	T K E	
GCTAAAGAAA AAGGTTTAGC	TCAAGTGGCC	CATCTCAGTA	AAGAAGAAGT	TGCGGCAGTC	600
A K E K G L A	Q V A	H L S K	E E V	A A V	
AATGAAGCAA AAAGACAAGG	ACGCTATACT	ACAGACGATG	GCTATATTTT	TAGTCCGACA	660
N E A K R Q G	R Y T	T D D G	Y I F	S P T	
GATATCATTG ATGATTTAGG	AGATGCTTAT	TTAGTACCTC	ATGGTAATCA	CTATCATTAT	720
D I I D D L G	D A Y	L V P H	G N H	Y H Y	
ATTCCTAAAA AGGATTTGTC	TCCAAGTGAG	CTAGCTGCTG	CACAAGCCTA	CTGGAGTCAA	780
I P K K D L S	P S E	L A A A	Q A Y	W S Q	
AAACAAGGTC GAGGTGCTAG	ACCGTCTGAT	TACCGCCCGA	CACCAGCCCC	AGGTCGTAGG	840
K Q G R G A R	P S D	Y R P T	P A P	G R R	
AAAGCCCCAA TTCCTGATGT	GACGCCTAAC	CCTGGACAAG	GTCATCAGCC	AGATAACGGT	900
K A P I P D V	T P N	P G Q G	H Q P	D N G	
GGCTATCATC CAGCGCCTCC	TAGGCCAAAT	GATGCGTCAC	AAAACAAACA	CCAAAGAGAT	960
G Y H P A P P	R P N	D A S Q	N K H	Q R D	
GAGTTTAAAG GAAAAACCTT	TAAGGAACTT	TTAGATCAAC	TACACCGTCT	TGATTTGAAA	1020
E F K G K T F	K E L	L D Q L	H R L	D L K	
TACCGTCATG TGGAAGAAGA	TGGGTTGATT	TTTGAACCGA	CTCAAGTGAT	CAAATCAAAC	1080
Y R H V E E D	G L I	F E P T	Q V I	K S N	
GCTTTTGGGT ATGTGGTGCC	TCATGGAGAT	CATTATCATA	TTATCCCAAG	AAGTCAGTTA	1140
A F G Y V V P	H G D	H Y H I	I P R	S Q L	
TCACCTCTTG AAATGGAATT	AGCAGATCGA	TACTTAGCTG	GCCAAACTGA	GGACAATGAC	1200
S P L E M E L	A D R	Y L A G	Q T E	D N D	
TCAGGTTCAG AGCACTCAAA	ACCATCAGAT	AAAGAAGTGA	CACATACCTT	TCTTGGTCAT	1260

S	G	S	E	Н	S	K	P	S	D	·K	Ė	V	Т	Н	·T	F	L	G	Н	
C(R	GCA1 I	rca <i>i</i> K	AAG A		ACGG G	SAAA K	AGG G	CTT L	'AGAT D	GG G	TAA K	ACC P	AT Y	ATGA D	TAC	CGAG S	TGA D	ATGC A	TTAT Y	1320
G' V	rtti F	TAC S	STA K	AAGA E	ATC S	CAT	TCA H	TTC	AGTG V	GA D	TAA K	ATC S	AG G	GAGT V	TAC T	CAGC A	TAP K	ACA H	CGGA G	1380
GZ D	ATC <i>E</i> H	ATTI F	CC H	ACTA Y	TAT I	'AGG G	ATT F	TGG G	AGAA E	CT L	TGA E	ACA Q	TA. Y	ATGA E	GTI L	GGA D	TGA E	AGGT V	CGCT A	1440
A. N	ACTO W	GGI V	GA K	AAGC A		AGG G	TCA Q	AGC A	TGAT D	GA E	GCT L	TGC A	TG A	CTGC A	TTI L	'GGA D	TCA Q	AGGA E	ACAA Q	1500
G(G	GCAA K	AGA E	AA K	AACC P	ACT L	CTT F	TGA D	CAC T	TAAA K	AA K		GAG S	TC R	GCAA K	AGT V	'AAC T	AAA K	AGA D	TGGT G	1560
A/ K	AAGI V	'GGG G	CT Y	ATAT M	'GAT M	GCC P	AAA K	AGA D	TGGT G	AA K	GGA D	CTA Y	TT F	TCTA'	TGC A	TCG R	TGA D	TCA Q	ACTT L	1620
G <i>I</i> D	TTT L	'GAC T	TC Q	AGAT I	TGC A	CTT F	TGC A	CGA E	ACAA Q	GA. E	ACT. L	AAT M	GC L	TTAA K	AGA D	TAA K	GAA K	GCA H	TTAC Y	1680
CO R	TTA Y	TGA D	CA I	TTGT V	TGA D	CAC T	AGG' G	TAT I	TGAG E		ACG. R	ACT L	TG A	CTGT: V	AGA D	TGT V	GTC S	AAG S	TCTG L	1740
CC P	GAT M	GCA H	TG A	CTGG G	TAA N	TGC A	TAC'	TTA Y	CGAT D	AC'	TGG G	AAG S	TT S	CGTT:	rgt V	TAT I	CCC	ACA H	TATT I	1800
G <i>P</i> D	ATCA H	TAT	CC H	ATGT V	CGT V	TCC P	GTA' Y	rtc: S	ATGG W	TTC L	GAC(T	GCG R	CG D	ATCA(GAT I	TGC A	AAC. T	AGT V	CAAG K	1860
TA Y	TGT V	GAT M	GC Q	AACA H	CCC P	CGA E	AGT'	rcg: R	rccg P	GA!	rgtz V	ATG W	GT S	CTAA(K	GCC. P	AGG G	GCA H	TGA E	AGAG E	1920
TC S	AGG G	TTC S	GG V	TCAT I	TCC. P	AAA N	TGT: V	raco T	GCCT P	CT!	rga: D	TAA K	AC R	GTGCT A	rgg G	TAT M	GCC.	AAA(N	CTGG W	1980
CA Q	I AA	TAT I	CC H	ATTC S	TGC' A	TGA E	AGA/ E	AGT: V	rcaa Q	AAA K	AGC(A	CCT.	AG A	CAGA <i>I</i> E	AGG' G	TCG R	TTT'	TGC A	AACA T	2040
CC P	AGA D	CGG G	CT Y	ATAT'	TTT(F	CGA D	TCC?	ACG <i>I</i> R	AGAT D		ĽŢŦ(L	GGC A	CA K	AAGAA E	AAC' T	TTT F	TGT: V	ATG(GAAA K	2100
GA D	TGG G	CTC S	CT F	TTAG S	CAT(CCC P	AAGA R	AGC! A	AGAT D	GG(G	CAGT S	rtc. S	AT L	TGAGA R	AAC T	CAT	TAA! N	TAA <i>I</i> K	ATCT S	2160
GA D	TCT: L	ATC S	CC Q	AAGC! A	TGA(E	gtg W	GCA/ Q	ACA <i>I</i> Q	AGCT A	CA <i>I</i> Q	AGA(E	GTTI L	AT L	TGGCA A		GAA K	AAA' N	TAC:	rggt G	2220
GA D	TGC' A	TAC'	TG D	ATAC:	GGA'	raa K	ACCO P	CAAA K	AGAA E	AAC K	GCAA Q	ACA(Q	GG A	CAGAT D		GAG S	CAA: N	rga <i>i</i> E	AAAC N	2280
CA Q	ACA(Q	GCC P	AA S	GTGAZ E	AGC(A	CAG S	TAAA K	AGAA E	AGAA E	AAA K	AGA <i>A</i> E	ATCA S	AG D	ATGAC D		TAT I	AGA(_	TTTA L	2340
CC P	AGA(CTA' Y	TG G	GTCTI L	AGA:	rag R	AGCA A	ACC T	CTA L				A7 I	TCAAT N	'CAI Q		AGC! A		AAAA K	2400
GC A	TAA! N	rat(CG D	ATCC:	raac K	GTA Y	TCTC L	ATT I	TTTC	CA.	ACC <i>P</i> P	AG A Z E	AG G	GTGTC V	CAI Q	ATT F	TTA:	raat N	TAAA K	2460

AATGGTGAAT N G E L	TGGTAACTTA V T Y	TGATATCAAG D I K	ACACTTCAAC T L Q Q	AAATAAACCC I N P	TTAACCAAAA	2520
GAAGATCTCA	TTGTTAAAGC	ACTGCTTTGT	CAAAGCAAGT	TACGGTGATT	TTGAAGTCAT	2580
TCTATGTAAC	GAGTAGTGAT	AAAAGTTGGA	TAATAGCGGT	TTTCTTTTGC	AAAGAAATGG	2640
TATCCATGTT	AGAATAGTAA	AAAAAGAGGA	GGATTCTTGG	ACTAATGTCA	AATAAGTAGA	2700
CAGAAAACTG	TGTTATTTTA .K	TTGCGTTAAA I A N F	ATAATTTTCT Y N E	TCTTTCTGAT E K Q	TAGGGGTTAG N P T L	2760
TCCTAGATTA	GCCGTATGTG	GGTTGTAATT	GTTATAAAAA	TTCTCAATGT	ATTCAAAGCA	2820
G L N	A T H	P N Y N	N Y F	N E I	Y E F C	
GTCTAATTGA	ACCTGTTTGA	TATTTTGATA	ATGTTTTCGG	TTGATTTGTC	TATGCTTTAA	2880
D L Q	V Q K	I N Q Y	H K R	N I Q	R H K L	
ATACTTGAAA	AATGCTTCAG	TTACGGCATT	ATCATAAGGA	TATCCAGGAT	TAGAAAAAGA	2940
Y K F	F A E	T V A N	D Y P	Y G P	N S F S	
ATGCATGATA H M <	TTGGCACTGC	ACCCTAATAG	TGAGACGCAA	GAAAAACACT	TTTAGGCAAT A I	3000
CAGTTTTCTG	TACTGTACAG	GCGACTGGTC	GTTTAATCTC	TGTTGAATTC	TAGTTTCATT	3060
L K R	Y Q V	P S Q D	N L R	Q Q I	R T E N	
ATAAAATGTA	ATGTAATTTT	TAACAATATT	TGTTATACTA	TCTTTGTTGT	ATTTTCTCCT	3120
Y F T	I Y N	K V I N	T I S	D K N	Y K R R	
ATTATGGAAA	TAAAAGGTTT	CAGTCTTTAG	GACGGTGTGA	AACCATTCAA	TACAGGCATT	3180
N H F	Y F T	E T K L	V T H	F W E	I C A N	
ATCTGCAGGT	GTTCCTTTTC	GAGACATTGA	GCGGATAATG	TCTTTTTCCG	TGCAAGCCTG	3240
D A P	T G K	R S M S	R I I	D K E	T C A Q	
GTAGTAAGCC Y Y A <	ATAGAAGTAT M	ACACTGAGCC	TTGGTCACTG	TGTAAGATTG	CTCCTTTATT	3300
TAGGCAATTT	TAACTGATTA	AGGGTGTCTA	GTACAAAATC	CGTGTCCTGA	CAATCTGAGA	3360
K P L K	L Q N	L T D	L V F D	T D Q	C D S	
TAGTGTAAGC	TATAATTTCT	CGGTTATAGA	GATTCATAAT	TGATGAGAGA	TACAATTTAC	3420
I T Y A	I I E	R N Y	L N M I	S S L	Y L K	
AGTTACCGAA	ATATAGGTAG	GTAATATCTG	TTACGAGCTT	TTCCTTAGGC	TTATCGGCAT	3480
C N G F	Y L Y	T I D	T V L K	E K P	K D A	
GGAAATCCCG H G D R	ACTCAATTTA S L K		AATAATAAGC L Y Y A		TTGGGAACTT N P V	3540
TCTTGGTACG	TGTCCGACAA	AGCCAGCCAT	TATTTTTCAT	GATACGATAG	ACTTTCTTTG	3600
K K T R	T R C	L W G	N N K M	I R Y	V K K	
TATTAACAGT	CAATCCGTGG	ATTTTTTTGA	GCAATCGTGT	AATGGTACGA	TAGCCATAAA	3660
T N V T	L G H	I K K	L L R T	I T R	Y G Y	
TAAAGTGATT I F H N	CTCCATACAG E M <	AGCTGTTCAA	TTAATTCAAT	AAGGTCATCT	TTTTTTGCGG	3720

CTTCTCATAC	TCCTTTTTCC	AACGGTAATA	GGTCGACCGC	TTGACCTTAA	AACAGTCTAG	3780
AATGAAAACT	ATCGGGTAGT	TGTTTTTATA	GTCTTCCACA	AGCTTGATAA		3840
ATCGATTTCC I S K	TTATCAAGCC R I L G		TTTAAGAGGT K L L	CAACCTGTAA D V Q L	-101111101	3900
TCCACTTCAG E V E	ACAGATGTTC S L H E		CCGTAGGTAT G Y T	ATTGCTTGCC Y Q K G	1-10110011011	3960
TGAAAACGAT H F R	AAAGCTCCTC Y L E E		CATTTCATCC W K M	AAGTATAGAT W T Y I	TTGACTATTA Q S N	4020
TTTTTGATGC N K I	CTAAAGTCTC G L T E		CTGTTAGACT R N S	TGCCTGCTTT K G A K	CTTCATATCG K M D	4080
ATGCAAGCCA I C A	GCTTAGTTTC L K T E		GCTTTTTTAA A K K	CCATAATAAA V M	ACATTCCTGT	4140
TTCTAGTTTA	CTAAATTTCA	ACAGGAGTGT	TTTTCTTTTG	TCTCATTTTA	GGGATTCAGT	4200
GCCTATTGTT	GTCATCAATT	ATTTTTCTAA	ATTCCCCGGA	CTTAAATTGT	GACCCTTGGT	4260
CGGAATGAAA	GAGAAGTGTT	CCTTCAATCT	TTCTTTTATT	AAGTGAAAAG	GCAACACTTT	4320
TCTGTACAAC	ATTTATAAAG	TGTTTTTCTA	GGCAATTAAT . A I L	CTTTTAGTCA R K T	TTGGTGTTTG M P T Q	4380
GTAGTTGAGA	CTACCATGAA	TGCGGTGGTA	ATTCCACCAA	TGAACATAGT	CTTTAGTCTT	4440
Y N L	S G H	I R H Y	N W W	H V Y	D K T K	
AAGAGCTAGT	TCTTCCAGCA	ATTGAAAGGT	TTCTTGATAA	ACAAATTCAA	TTTTGAAAGC	4500
L A L	E E L	L Q F T	E Q Y	V F E	I K F A	
ACGATACGTA	CTTTCAGCTA	CGGCATTGTC	ATAAGGATAA	CCAGCCTGAC	TAAGCGAACG	4560
R Y T	S E A	V A N D	Y P Y	G A Q	S L S R	
TGTGATTCCA	AAGGCTTCCA	ATATTTCATC	AATTAACTGA	TTATCAAACT	CTTTGCCACG	4620
T I G	F A E	L I E D	I L Q	N D F	E K G R	
ATCTGAATGG	AACATCTTGA	CTTTGGTCAG	GGCGTAAGGG	ATGCTTTGTA	TGGCTTGCTT	4680
D S H	F M K	V K T L	A Y P	I S Q	I A Q K	
AACGAGTTCA	GCGGTCTTGT	GCCAACCAAG	AGACAGGCCG	ATGATTTCAC	GGTTGTATAG	4740
V L E	A T K	H W G L	S L G	I I E	R N Y L	
GTCAATGATG	AGGCAAACAT	AAGCCCAACG	ATTGCCTACA	CGAACATAGG	TTAAGTCAGT	4800
D I I	L C V	Y A W R	N G V	R V Y	T L D T	
GACTAAGGCT	TGTAGTGGTC	TTTCTTGCTT	AAATTGCCTG	TCTAAGTGGT	TGGGAATAGG	4860
V L A	Q L P	R E Q K	F Q R	D L H	N P I P	
GGCTTCATTC	TTGCCTCTAG	AATGTGGTTT	GAAGGTGGCT	TTCTGATAAA	CAGAAACCAA	4920
A E N	K G R	S H P K	F T A	K Q Y	V S V L	
ATTGAGTCGC	TTCATAATGC	GTCGAATCCG	ACGACGTGAA	AGTGTGATAC	CTTCGTTATT	4980
N L R	K M I	R R I R	R R S	L T I	G E N N	
CAAGCATATT	TTGATTTTTC	TGGATCCGTA	TCTAGACTCG	CTATCGAGAA	AAATTCTTTT	5040
L C I	K I K	R S G Y	R S E	S D L	F I R K	

AATAGTTTCT TCAAACTCCG TTTCAGATAC TGACTCCACG GCTTGATAGT AATAACTTGA 5100

		E S V S I		AGT AATAACTTGA Y Y S S	5100
		ACATCTT TGAAAT C M K S 3			5160
GATTATTTCC CT I I E I (SEQ ID NO	R K T G	AATCACC GCTGCT Y D G S S FIG. 3	S A K P Y		5215
FGSALSTVEV (SEQ ID NO		LYRLSCCHFT FIG. 3			40
MGIÄTKDNOT	AVIDDSKCKA	WA DWWNIWMAN	OTCAUDCTCA		
		KAPKTNKTMD AIISEELLMT			50
				VINEILDGYV	100
	LKPGSKRKNI	22-00-2		GLAQVAHLSK	150
				GNHYHYIPKK	200
		GARPSDYRPT			250
HQPDNGGYHP	APPRPNDASQ	NKHQRDEFKG	KTFKELLDQL	HRLDLKYRHV	300
EEDGLIFEPT	QVIKSNAFGY	VVPHGDHYHI	IPRSQLSPLE	MELADRYLAG	350
QTEDNDSGSE	HSKPSDKEVT	HTFLGHRIKA	YGKGLDGKPY	DTSDAYVFSK	400
ESIHSVDKSG	VTAKHGDHFH	YIGFGELEQY	ELDEVANWVK	AKGQADELAA	450
ALDQEQGKEK	PLFDTKKVSR	KVTKDGKVGY	MMPKDGKDYF	YARDQLDLTQ	500
		VDTGIEPRLA			550
		QIATVKYVMQ			600
		SAEEVQKALA			650
		RTINKSDLSQ			700
		EASKEEKESD			750
		VQFYNKNGEL			
		· Xr THIMADTT	ATTUTUTHOO	TME	793

FIG. 3c

(SEQ ID NO:15)

MTDPNYRFKQ	SDVINEILDG	YVIKVNGNYY	VYLKPGSKRK	NIRTKQQIAE	5(
QVAKGTKEAK	EKGLAQVAHL	SKEEVAAVNE	AKRQGRYTTD	DGYIFSPTDI	100								
IDDLGDAYLV	PHGNHYHYIP	KKDLSPSELA	AAQAYWSQKQ	GRGARPSDYR	150								
PTPAPGRRKA	PIPDVTPNPG	QGHQPDNGGY	HPAPPRPNDA	SQNKHQRDEF	200								
KGKTFKELLD	QLHRLDLKYR	HVEEDGLIFE	PTQVIKSNAF	GYVVPHGDHY	250								
HIIPRSQLSP	LEMELADRYL	AGQTEDNDSG	SEHSKPSDKE	VTHTFLGHRI	300								
KAYGKGLDGK	PYDTSDAYVF	SKESIHSVDK	SGVTAKHGDH	FHYIGFGELE	350								
QYELDEVANW	VKAKGQADEL	AAALDQEQGK	EKPLFDTKKV	SRKVTKDGKV	400								
GYMMPKDGKD	YFYARDQLDL	TQIAFAEQEL	MLKDKKHYRY	DIVDTGIEPR	450								
LAVDVSSLPM	HAGNATYDTG	SSFVIPHIDH	IHVVPYSWLT	RDQIATVKYV	500								
MQHPEVRPDV	WSKPGHEESG	SVIPNVTPLD	KRAGMPNWQI	IHSAEEVQKA	5 5 0								
LAEGRFATPD	GYIFDPRDVL	AKETFVWKDG	SFSIPRADGS	SLRTINKSDL	600								
SQAEWQQAQE	LLAKKNTGDA	TDTDKPKEKQ	QADKSNENQQ	PSEASKEEKE	650								
SDDFIDSLPD	YGLDRATLED	HINQLAQKAN	IDPKYLIFQP	EGVQFYNKNG	700								
ELVTYDIKTL	QQINP (SEC	ID NO:16)			715								
FIG. 3d													
MHSFSNPGYP	YDNAVTEAFF	KYLKHRQINR	KHYQNIKQVQ	LDCFEYIENF	50								
YNNYNPHTAN	LGLTPNQKEE	NYFNAIK (S	SEQ ID NO:17	')	77								
		FIG. 3	е										
MAYYQACTEK	DIIRSMSRKG	TPADNACIEW	FHTVLKTETF	YFHNRRKYNK	50								
				Q ID NO:18)	86								
		FIG. 3	İ										
MENHFIYGYR	TITRLLKKIH	GLTVNTKKVY	RIMKNNGWLC	RTRTKKVPNL	50								
				SSTMNT VNDE									

FIG. 3g

126

IIAYTISDCQ DTDFVLDTLN QLKLPK (SEQ ID NO:19)

MVKKAYSWET	KLACIDMKKA	GKSNRVIMET	LGIKNNSQIY	TWMKWYENEE	50
LYRFHQGVGK	QYTYGKGLEH	LSEVEQLQLQ	VDLLKKYRGL	IRKSIK	96
(SEQ ID NO:	20)				

FIG. 3h

IRYPKASSGD	YGTKREIITA	NKDKYSISKM	CRWLNMPHSS	YYYQAVESVS	50
ETEFEETIKR	IFLDSESRYG	SRKIKICLNN	EGITLSRRRI	RRIMKRLNLV	100
SVYQKATFKP	HSRGKNEAPI	PNHLDRQFKQ	ERPLQALVTD	LTYVRVGNRW	150
AYVCLIIDLY	NREIIGLSLG	WHKTAELVKQ	AIQSIPYALT	KVKMFHSDRG	200
KEFDNQLIDE	ILEAFGITRS	LSQAGYPYDN	AVAESTYRAF	KIEFVYQETF	250
QLLEELALKT	KDYVHWWNYH	RIHGSLNYQT	PMTKRLIA (S	SEO ID NO:21	288

FIG. 3i

N	TTT(L >	GAA K	AG A	CAGA E	ATT: L	ATC S	TGT: V	AGAZ E	AGAT D		GCA Q		TA T	CAGC:	AAC. T	AGT V	TTA: Y		raaa K	60
TC' S	IGC' A	TCA'	rg G	GTTC:	AACI T	ACC P	ACA: Q	AGAZ E			raa: N			CGAC'		TTT L	AGC:		TAT Y	120
CT L	AAG' S	TCAZ Q	AT F	TTGA' D	TTT'	rga E	AGG'	rcc: P	IGCT A		TGC' A			TAGA' D	TGT' V	TAC T	AGC(CATT	180
AT'	rca H	CGAZ E	AG D	ACTT(CTC S	AGG G	TGAZ E	AAA <i>I</i> K	ACTT L		AGT <i>I</i> V			ATGA:	AGA' D	IGA D	CTG1 C	CATO M	GGA G	240
CCI P	ATT(L	GAG(S	CA M	TGAA' N	rgc <i>i</i> A	AGG G	TGT	CTT(F	CCAG Q	TT:	rga: D		AA T	CTAA'	rga: D	rga D	TAAT N		TATC I	300
GC'. A	rct: L	raat N	rT F	TCCG'	Y Y	CCC P	ACAI Q	AGG(G	GACA T		rgc: A			CTATO	CCAI Q	AAC	TAAC		rgag E	360
AAI K	ACT:	raac N	CG G	GAGT: V	rga <i>i</i> E	AAA K	AGT(GACT T	CTT L	TC: S		CCA!	rg E	AACA(CAC T	ACC P	ACAC H	TAT Y	rgta V	420
CC: P	TAT(M	GGA(D	D D	ATGAI E	ATT <i>I</i> L	AGT V	ATCA S	AAC(T	CTTA L		AGC: A			ATGA! E	AAA(K			GGI G	CTT L	480
AA/ K	AGG <i>I</i> G	ACAT H	rg E	AACA(GGTT V	TAT I	TGG:	rgg1 G	r G GG G		ATT:		rc R	GCTT!		rga E		GG1 G		540
GC <i>I</i> A	ATA(Y	CGG1 G	PT A	CCAT(GTT(F	CCC P	AGG! G	AGAT D	rgaa E	AA(N	CACT T	TAT(M	GC H	ATCA/ Q	AGC: A			STAC Y		600
CCI	CTTA	AGA	A.A	TATA	TTT	CCG	TTC	GC1	GCT	ATO	CTAC	CGC	AG	AAGC	rat(CTA	TGAA	TTF	ATC	660

E	PI	E	N	I	F	R	S	A	A	I	Y	A	E	A	I	Y	E	L	I	
F	AAAT	'AAA	ATA	ATC	CTT	AAAC	TAA	AT <i>F</i>	ATGTG	ATO	CAA	TGF	ATA	AAGG	GTG	GTG	AAG	ACA	TGAA	720
F	AGTG	TCT	TTG	CCI	CTT	TTCA	TAA	AGGI	TAGA	TTI	rgg.	AGA	CT	TTAT M !_	GAC T	CTGA D	_	GGA E	AAAA K	780
Į	ATTA I	TTA K	AAG A	CAA	TAA K	AAAG S	TGA D	TTC S	CACAG Q	AAT N	CA Q	AAA N	TT. Y	ATAC T	AGA E	AAA N	TGG G	TAT I	TGAT D	840
E	CTT	TGT F	TTG A	CTG A	CTC P	CTAA K	AAC T	AGC A	TAGG R	ATC I				TTGG G	CCA Q	AGC A	ACC P	TGG G	TTTA L	900
K	AAA T	CTC Q	AAG E		CAA R	GACT L	CTA Y	TTG W	GAAA K		'AA K			GAGA D	TCG R		ACG R	CCA Q	GTGG W	960
C	TTG G	GAG V	TTG D	ATG E	AAG E	AGAC T	ATT F	TTA Y	CCAT H	TCT S				TTGC A		'TTT L	ACC'	_	AGAT D	1020
Ŀ	Y	Y	P	G	K	G	K	S	G	D	L	P	P	CTAG R	K	G	F	A	E	1080
A	TAA. W	GGC. H	ACC P	CTC L	TTA' I	TTTT L	AAA K	AGA E	AATG M	CCT P	'AA' N	rgt V	TC Q	AATT L		CTT L	GCT/ L		TGGT G	1140
Q	AGT. Y	ATG A	CTC Q	AGA K	AAT. Y	ATTA Y	TCT L	TGG G	AAGC S	TCC				AAAA N	TCT L	AAC T	AGAZ E	AAC T	AGTT V	1200
A	AAG A	CTT. Y	ACA K	AAG D	ACT: Y	ATCT L	ACC P	CGA D		TTA L		CCT L	GG V	TTCA	CCC P	ATC S	ACC	GCG2 R	AAAT N	1260
Q	AAA I	TTT W	GGC L	TAA K	AGA K	AGAA N	TCC.	ATG W	GTTT F	GAA E	AA. K	AGA D	TC L	TAAT		TGA D	TTTZ L	ACAZ Q	AAAG K	1320
A	TAG' V	TAG A	CAG D	ATA I	TTT' L	TAAA K	AGA D		AGGA	TAG	GAC	STT	GG	TATG	AGA R				CTAC L H	1380
A	CAC	GTA' Y	PTT F	TTC S	CTA' Y	TGAT D	TGT	CAA Q	ACGG T A	CAT		GAG E		CTAT			GGT:		ACAG I G	1440
G	ŤGAZ E	ATT' F	rat I	CAC	GAC T	AGAA E	CAT H		GATT D L		CAA N			TTAC			CAA(GATG V	1500
T	TCC' P	rga: D	TTA Y	TAG' S	TGC: A	TAT Y	TGT	CAA Q	AAAA K I	TAG D		CAT		TAAT(AAA K	TATO		AATC N R	1560
G.	ATT!	raaz K	AAA K	AGG:	AAT! I	rgaa E	ATC	GGT G	TATT Y F	TTA K	AA (SAT.	AG R	GGAA'			ATTT I I		GATT O Y	1620
A	TTTI L	AAAI K	AAA N	TAA	AGA! E	ATTT F	GAT'	TTA L	AAAC K L	TAT L				CCAT(AAT N			TATG Y D	1680
A	TTA: Y	ICT(GCA Q	AGA E	AGA! E	AGCT A	CTG.	AAA K	GTAC V P		CAF	***	GG G	AGCT:		AGC S	AGAT R I			1740
A	ATC	GTA:	rgg	AAT'	TTG	CCAT	AGG	CCG	TGTG	GAA	.GCG	SCA	CG	TTTT	AGC	TCA	CTTI	'GA	TAT	1800
G	GTT	rtc	GTA	AGT'	TAAI	ACTT	AGA'	TGT.	AGAA	GAT	TTA	AA	AC	CGTT	rga.	AAC	GCA	TTC	SAAG	1860
C	GCA:	rtt:	rca	TAA	AGA:	rgtt	ATC'	TAA	GGGG	TTA	GCI	TT	TG	AACT	AAA	TAC	CAAA	\TCC	بالبلات	1920

TATCTATATG	GGAATGAAAA	ACTTTATCGC	TATGCTTTAG	AGATACTCAA	ACAGCTTGGT	1980
TGTAAACAAT	ACTCTATAGG	CTCTGACGGT	CATATTCCTG	AACATTTTTG	TTATGAATTT	2040
GATAGACTTC	AAGGTCTGCT	AAAGGACTAT	CAAATTGATG	AAAATCATTT	GATATGAGGA	2100
AATTTTTGAT	AAAAAAGCTA	GGCAATATTG	CTTAGCTTTT	TTGTAATGCT	ATTGATAGTT	2160
TTAGTGAAAA	TTTCAAAAAA	ATAAAGAAAT	CATTTACTTG	TTGCAAGCGC	TTGCGTAAAT	2220
TGTTATGATT	TTATTGGTAA	CAATTCATTA	AAAAAGGAGA		AAGAAAAGAC R K D	2280
TTATTTGGTG L F G D	ATAAACAAAC K Q T	TCAATACACG Q Y T	ATTAGAAAGT I R K L		AGTAGCTTCA V A S	2340
GTTACAACAG V T T G	GGGTATGTAT V C I	TTTTCTTCAT F L H	AGTCCACAGG S P Q V	TATTTGCTGA F A E	AGAAGTAAGT E V S	2400
GTTTCTCCTG V S P A	CAACTACAGC T T A	GATTGCAGAG I A E	TCGAATATTA S N I N	ATCAGGTTGA Q V D	CAACCAACAA N Q Q	2460
TCTACTAATT S T N L	TAAAAGATGA K D D	CATAAACTCA I N S	AACTCTGAGA N S E T	CGGTTGTGAC V V T	ACCCTCAGAT P S D	2520
ATGCCGGATA M P D T	CCAAGCAATT K Q L	AGTATCAGAT V S D	GAAACTGACA E T D T	CTCAAAAGGG Q K G	AGTGACAGAG V T E	2580
CCGGATAAGG P D K A	CGACAAGCCT T S L	GCTTGAAGAA L E E	AATAAAGGTC N K G P		TAAAAATACC K N T	2640
TTAGATTTAA L D L K	AAGTAGCACC V A P	ATCTACATTG S T L	CAAAATACTC Q N T P		TTCTCAAGCT S Q A	2700
ATAGGTGCTC I G A P	CAAGCCCTAC S P T	CTTGAAAGTA L K V	GCTAATCAAG A N Q A	CTCCACGGAT P R I	TGAAAATGGT E N G	2760
TACTTTAGGC Y F R L	TACATCTTAA H L K	AGAATTGCCT E L P	CAAGGTCATC Q G H P	CTGTAGAAAG V E S	CACTGGACTT T G L	2820
TGGATATGGG W I W G	GAGATGTTGA D V D	TCAACCGTCT Q P S	AGTAATTGGC S N W P	CAAATGGTGC N G A	TATCCCTATG I P M	2880
ACTGATGCTA . T D A K	AGAAAGATGA K D D	TTACGGTTAT Y G Y	TATGTTGATT Y V D F	TTAAATTATC K L S	TGAAAAACAA E K Q	2940
CGAAAACAAA R K Q I	TATCTTTTTT S F L	AATTAATAAC I N N	AAAGCAGGGA K A G T	CAAATTTAAG N L S	CGGCGATCAT G D H	3000
CATATTCCAT '	TATTACGACC	TGAGATGAAC	CAACTTTCCA	ТТСТПСТТТ		3060
	AACCCCTCAA.	AGAAGGGTAT	GTCCGTATTA	ACTATTTGAG '	TTCCTCTAGT S S S	3120
AACTATGACC I N Y D H	ACTTATCAGC A	ATGGCTCTTT W L F	AAAGATGTTG (K D V A	CAACCCCYTC 1	AACAACTTGG T T W	3180
CCAGATGGTA (P D G S	GTAATTTTGT (N F V	GAATCAAGGA N Q G	CTATATGGAA (L Y G R	GGTATATTGA :	IGTATCACTA V S L	3240

K	T	N N	A.	. CCAA	E E	I I		F	TCTA L	I	CTT L	'AGA D	TG E		TAA K	GAC T	AGG G	AGA D	ATGCA A	3300
G1 V	GAA K	AGT V	TC	AACC P	CAA N	CGA D	CTA Y	TGT V	TTTT F			TTT L		CTAA N	CCA H	TAA N	CCA Q	AAT I	TTTT F	3360
GI V	'AAA K	AGA D	TA K	AGGA D	TCC P	AAA K	GGT V	TTA Y	TAAT N			TTA Y		ACAT I		TCA Q	AGT V		GCTA L	3420
AA K	AGGA D	ATGC A	CC Q	AACA Q	AAT I	TGA D	TTT L	AAC T	AAGT S			AGC A		GTTT F		AAC T	TCT L	_	TGGG G	3480
GT V	'AGA D	TAA K	AA T	CTGA E	AAT I	TTT L	AAA K	AGA E	ATTG L			GAC T		ATAA K		TCA Q	AAA N	TGC A		3540
CA Q	AAT I	TTC S	TG D	ATAT	CAC T	TCT L	CGA D		TAGT S			TCT L		TAAT		CAA K	AGG G		CTTT F	3600
AA N	TCC P	TAA K	AC Q	AAGG G	TCA H	TTT F	CAA N	CAT. I	ATCT S	TA: Y	ΓΑΑ' Ν	TGG' G	TA N	ACAA: N	rgt V	CAT M	GAC.	AAG R	GCAA Q	3660
TC S	TTG W	GGA. E	AT F	TTAA K	AGA D	CCA Q	ACT'	TTA Y	TGCT A	TA: Y	rag' S	rgg/ G	AA N	ATTTA L	AGG" G	IGC A	AGT V	_	CAAT N	3720
CA Q	AGA D	TGG' G	TT S	CAAA K	AGT' V	TGA E	AGC(CAG S	CCTC L	TG(W	STC2	ACC(GA S	GTGCT A	rga: D	rag s	TGT V	CAC' T	TATG M	3780
AT I	TAT I	TTA' Y	rg D	ACAA K	AGA' D	TAA N	CCA Q	AAA N	CAGG R	GTT V	rgt <i>i</i> V	AGC(A	GA T	CTACO	CCC P	CCT L	TGT	GAAZ K	AAAT N	3840
AA N	TAA K	AGG: G	rg V	TTTG(W	GCA(Q	GAC T	GATI I	ACT' L	TGAT D	ACT T		ATT <i>I</i> L	AG G	GTATT I	TAAZ K	AAA N	CTA:	TAC'	rggt G	3900
TA Y	CTA' Y	TTA: Y	rc L	TTTA(Y	CGAZ E	AAT I	AAA! K		AGGT G	AAC K	GAT D	raac K	G V	TTAAG K	ATT I	rtt L	AGA:	rcc: P	TAT Y	3960
GC. A	AAA(K	GTC <i>I</i> S	T. L	TAGCA A	AGA(E	gtg W	GGA:	rag: s	TAAT N	ACI T	GTI V	raat N	D D	ATGAT D	TATI	TAA K	AAC(raaa K	4020
GCI A	AGC: A	TTTT F	V V	TAAAT	rcc <i>i</i> P	AAG S	TCAA Q		rgga [.] G		CA; Q		TT L	TAAGI S	TTI F	GC A	TAA <i>I</i> K	AAT1 I	GCT A	4080
AA' N	rtt: F	raa <i>r</i> K	AG G	GAAG <i>I</i> R	ACA <i>I</i> Q	AGA D	TGCT A	rgt: V	I I	TAC Y	GA <i>F</i> E		AC H	ATGTA V	AGA R	AGA D	CTT(F	CACI T	TTCT S	4140
GA: D		ATCI S	L	TGGAT D	rgg <i>r</i> G	AAA K	ATTA L	AA. K	AAAT N	CAA Q	TTI F	G G	A. T	CCTTT F	'GCA A	AGC A	CTT1 F	TCA S	AGAG E	4200
AAI K	ACT <i>I</i> L	AGAI D	Y	ATTTA L	ACA(Q	GAA K	ATTA L	AGG <i>I</i> G	AGTT V	ACA T	CAC H	ATI I	C Q	AGCTT L	TTA L	CC P	GGT <i>I</i> V	ATTO L	SAGT S	4260
TA: Y	rtti F	rati Y	'G V	TTAAT N	GA <i>A</i> E	TA/ M	GGA1 D	'AAC K	STCA S	CGC R	TCA S	ACA T	AG A	CTTAC Y	ACT T	TC S	CTCF S	AGAC D	CAAT N	4320
AA' N	Y Y	CAAT N	T W	GGGGC G	TAT Y	GA D	CCCA	Q Q	SAGC S	TAT Y	TTI F	'GC'I A	C L	TTTCT S	GGG G	AT M	GTAI Y			4380
AA! K	ACC <i>P</i> P	AAAA K	G D	ATCCA P	ATCA S	AGC A	ACGI R	'ATC	GCC A	GAA E	TTA L	AAA K	C .	AATTA L	ATA I	CA H	TGAI	ATT T	CAT	4440

AAACGTGGCA K R G M		ACTTGATGTC L D V	GTCTATAATC V Y N H	ACACTGCAAA T A K	AACTTATCTC T Y L	4500
TTTGAGGATA F E D I	TAGAACCTAA E P N	TTATTATCAC Y Y H	TTTATGAATG F M N E	AAGATGGTTC D G S	ACCAAGAGAA P R E	4560
AGTTTTGGAG S F G G		AGGAACCACT G T T	CATGCAATGA H A M S	GTCGTCGTGT R R V	TTTGGTTGAT L V D	4620
TCCATTAAAT S I K Y	ATCTTACAAG L T S	TGAATTTAAA E F K	GTTGATGGTT V D G F	TCCGTTTTGA R F D	TATGATGGGA M M G	4680
GATCATGATG D H D A	CGGCTGCGAT A A I	TGAATTAGCT C	TATAAAGAAG Y K E A	CTAAAGCTAT K A I	TAATCCTAAT N P N	4740
ATGATTATGA M I M I	TTGGTGAGGG G E G	CTGGAGAACA '	TTCCAAGGCG F Q G D	ATCAAGGTCA Q G Q	GCCGGTTAAA P V K	4800
CCAGCTGACC P A D Q	AAGATTGGAT D W M	GAAGTCAACC (GATACAGTTG D T V G	GCGTCTTTTC V F S	AGATGATATT D D I	4860
CGTAATAGCT R N S L	TGAAATCTGG K S G	TTTTCCAAAT (GAAGGTACTC E G T P	CAGCTTTCAT A F I	CACAGGTGGC T G G	4920
CCACAATCTT P Q S L	TACAAGGTAT Q G I	TTTTAAAAAT A	ATCAAAGCAC I K A Q	AACCTGGGAA P G N	TTTTGAAGCA F E A	4980
GATTCGCCAG D S P G	GAGATGTGGT D V V	GCAGTATATT (ATAACCTTAC N L T		5040
GTGATTGCAA V I A K		(SEQ ID NO:	:22)			5058
	·	F	IG. 4a			

NLKAELSVED	EQYTATVYGK	SAHGSTPQEG	VNGATYLALY	LSQFDFEGPA	50
RAFLDVTANI	IHEDFSGEKL	GVAYEDDCMG	PLSMNAGVFQ	FDETNDDNTI	100
ALNFRYPQGT	DAKTIQTKLE	KLNGVEKVTL	SDHEHTPHYV	PMDDELVSTL	150
LAVYEKQTGL	KGHEQVIGGG	TFGRLLERGV	AYGAMFPGDE	NTMHQANEYM	200
PLENIFRSAA	IYAEAIYELI	K (SEQ ID	NO:23)		221

FIG. 4b

MTDLEKIIKA	IKSDSQNQNY	TENGIDPLFA	APKTARINIV	GQAPGLKTQE	50			
ARLYWKDKSG	DRLRQWLGVD	EETFYHSGKF	AVLPLDFYYP	GKGKSGDLPP	100			
RKGFAEKWHP	LILKEMPNVQ	LTLLVGQYAQ	KYYLGSSAHK	NLTETVKAYK	150			
DYLPDYLPLV	HPSPRNQIWL	KKNPWFEKDL	IVDLQKIVAD	ILKD	194			
(SEQ ID NO:24)								

FIG. 4c

MRDNHLHTYF SYDCQ	TAFED YINGFT	SEFI TTEHFDLSNI	YTGQDDVPDY	50
SAYCQKIDYL NQKYG	NRFKK GIEIGY	KDR ESDILDYLKN	KEFDLKLLSI	100
HHNGRYDYLQ EEALK	VPTKG AFSRLL	(SEQ ID NO:25	5)	126

FIG. 4d

MKRKDLFGDK	QTQYTIRKLS	VGVASVTTGV	CIFLHSPQVF	AEEVSVSPAT	50
TAIAESNINQ	VDNQQSTNLK	DDINSNSETV	VTPSDMPDTK	QLVSDETDTQ	100
KGVTEPDKAT	SLLEENKGPV	SDKNTLDLKV	APSTLQNTPD	KTSQAIGAPS	150
PTLKVANQAP	RIENGYFRLH	LKELPQGHPV	ESTGLWIWGD	VDQPSSNWPN	200
GAIPMTDAKK	DDYGYYVDFK	LSEKQRKQIS	FLINNKAGTN	LSGDHHIPLL	250
RPEMNQVWID	EKYGIHTYQP.	LKEGYVRINY	LSSSSNYDHL	SAWLFKDVAT	300
PSTTWPDGSN	FVNQGLYGRY	IDVSLKTNAK	EIGFLILDES	KTGDAVKVQP	350
NDYVFRDLAN	HNQIFVKDKD	PKVYNNPYYI	DQVQLKDAQQ	IDLTSIQASF	400
TTLDGVDKTE	ILKELKVTDK	NQNAIQISDI	TLDTSKSLLI	IKGDFNPKQG	450
HFNISYNGNN	VMTRQSWEFK	DQLYAYSGNL	GAVLNQDGSK	VEASLWSPSA	500
DSVTMIIYDK	DNQNRVVATT	PLVKNNKGVW	QTILDTKLGI	KNYTGYYYLY	550
EIKRGKDKVK	ILDPYAKSLA	EWDSNTVNDD	IKTAKAAFVN	PSQLGPQNLS	600
FAKIANFKGR	QDAVIYEAHV	RDFTSDRSLD	GKLKNQFGTF	AAFSEKLDYL	650
QKLGVTHIQL	LPVLSYFYVN	EMDKSRSTAY	TSSDNNYNWG	YDPQSYFALS	700
GMYSEKPKDP	SARIAELKQL	IHDIHKRGMG	VILDVVYNHT	AKTYLFEDIE	750
			RVLVDSIKYL		800
			EGWRTFQGDQ		850
			FITGGPQSLQ		900
GNFEADSPGD	VVQYIAAHDN	LTLHDVIAKS	I (SEQ ID	NO:26)	931

FIG. 4e

1 Q S	TTGACAGAA L T E	G GTCAACTTCO G Q L R	G TTCTGATATO S D I	CCTGAGTTCO P E F E	C GTGCTGGTGA R A G D	60
TACTGTACG	F GTTCACGCT	A AAGTTGTTGA K V V E	A AGGTACTCGC G T R		C AGATCTTTGA	120
AGGTGTTGT:	I ATCTCACGT	A AAGGTCAAGG K G Q G	AATCTCAGAA I S E	ATGTACACAC M Y T V	TACGTAAAAT	180
TTCTGGTGGT	T ATCGGTGTA	G AGCGTACATT E R T F	CCCAATTCAC	C ACTCCTCGTG	TTGATAAAAT D K I	240
CGAAGTTGTT E V V	CGTTATGGT	A AAGTACGTCG K V R R		TACTACTTAC	GCGCATTGCA A L O	300
AGGTAAAGCT	GCACGTATT	A AAGAAATCCG	TCGTTAATTT	TGATGATCAG	ATTTTAAAAA	360
TGCTTGGTTG	TTTGAGGAT	A GTAACTATGT	TTTAAAACTG	GACAACCAAG	ACGTAAAAA	420
TCTGCCTGTG	GGCAGTTTT:	T TTACTAGGTC	CCCTTAGTTC	AATGGATATA . H I Y		480
CCTAAGGAGT G L S	AATTGCTGGT Y N S T	T TCGATTCCGG	CAGGGGACAT C P V	ATTCATTGCA Y E N C		540
GGTTTAGAGO P K S	TATTTTGCCC	CAAATTTCTC L N R	TGATTAAGTT Q N L	TATCGTTCCT K D N R		600
TCTTGTAATT E Q L	GATGTGCGTA Q H A Y	A AACTTCTAAA ' V E L	GTGATATTTA T I N	AATTCTCGTG L N E H	ATCTAAAACT D L V	660
TGAGAGATGG Q S I	AAATTAGATA S I L Y	GCTTGCAAAT S A F	GTATGCCTGA T H R	GAGAGTGCAC L S H V	TCGTACCTCG R V E	720
CGACCAGTTA R G T	TTTTTCGGAT	AGTTTTATTG T K N	ACTGCATTAT V A N	TTGAAAGTTT N S L K	GTCGAATAAT D F L	780
CTGTCGTTTT R D N	TATTTTTTGT K N K T	AAATTCATGC F E H	AAAAAAAATA L F F	ATGTATCATT L T D N	GTCAATTGGT D I P	840
ATATTTCTGA I N R	TACTACTTTT I S S K	GTTTTTTGTT N K T	GGCAGGTATC P L Y	TTTGGTTGAA R Q N F	ATGATAATCC H Y D	900
CAAGTTTTAT W T K	TAATTGATAA N I S L	ATATTTGTTA Y K N	GTGTAATCAA T Y D	TATCATTAAC I D N V	TGTTAAACCT T L G	960
AAACATTCAG L C E	CGAAGCGCAT A F R M	GCCAGTTTTA	GCGATGAGGT	ATAACGCTGC	ATACGATTGA	1020
TGTTGTGATT	,	AATTTTTATC	AAGCGTAAGT	ATTCATTGGT	TTCAAGAAAT	1080
TTTATCTCTA	TTTACGCCCC	TTATTTTTTG	CTTTAACCTT	AGTGAATAAA	CAAAAATTTT	1140
TTTCTATATA	TCCCTCGTGA	ACAGCCATGG	ATACGCAGGC	TTTTACATGT	ATGTTAAAAC	1200
GCTTTACTGT	ATCTTGCACA	TGCGTTTGAC	TATAATGATT	TATGACTTGT	TGATATTTAG	1260

CCCACCCGTT GTCGCGTTTA CGGAAATACG CCATTGATAT ACTCCACATT AGCTAAAGAA 138 CAGGGTGTTC AAGGCTACCT TGATGGAAAA GGCTCTCTTA GAGATATTTG TAAATGGTAT 144 GATATCTCAA GTCGCTCTGT TCTCCAAAAG TGGATAAAAC GGTATACTAG TGGTGAAGAC 150 TTGAAAGCCA CTAGTAGAGG ATATAGCCGT ATGAAACAAG GAAGGCAAGC CACATTTGAA 156 GAACGTGTAG AGATTGTTAA CTACACCATT GCCCATGGGA AAGACTATCA AGCAGCTATT 162 GAGAAGTTTG GTGTTCCTA CCAACAAATT TATTCTTGGG TGCGTAAGCT TGAGAAGAAT 168 GGCTCACAAG GTTTGGTTGA TAGACCATT TATTCTTGGG TGCGTAAGCT TGAGAAGAAT 168 GGCTCACAAG GTTTGGTTGA TAGACCATG AAAGGGTTGG AAGGTAAGCC TGATTTAACC 174 GAGATTGAGC AACTTAACT CAAGATTAAA CAATTGGAGG AACGTAATCG TCTCTTAGAA 180 ATCGAGGTTA GTTTACTAAA AAAGTTAGAA GACATCAAAC GAGGAAACAG ACGGTAAGAC 186 TAGGTAAGCA TTTAGCGGAG TTCCAAGTAA TCAAGAATTA TTACGATGAG GAATCTAATG 192 TGCCTATTCA GGCCTTATGC CAACTCTTGA AGGGGTCTCG TTCAGGCTAT TACAAGTGGC 198 TCAATCGTCA AAAAACAGAT TTTGAGACAA AAAATACAAA GCTAATGGCT AAAAATCAAGG 204 AACTTCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAATC 206 GTCAACTTGG GACAACTTAA AACAAGAAAC GGATTCGTCG TATGACAACA TTTATTAATC 210 GTCAACTTGG GACAACTTAA AACAAGAAAC GGATTCGTCG TATGACAACA TTTTTTAATC 222 AAGAAAAATAT TCTTAATCGT GTTAGCCATG CTTGTACAAA AGCTGGTGAC AGATTTTACG 222 AAGAAAAATTA TCTTAATCGT GAATTTACAG CCACAGCTCA TAACCAGAAA TGGTGCACA 234 CTTATTCACTT TCTTCAATAC GGTCTGGGGG CAAAGCTTA TCTCAGTGCG ATTAAAGACC 234 CTGATTAACGG TTCTTATTC GCTTATGAGA TTAGTCACAA CAATGAAAT CACTTGTTAT 2400 GAAGACCATT AAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCTATCA TCCATAGCGA 246 CTTATCCATG TCCTCATATCC CAAAGAATA TCCAGGAGCC ACACCTATCA TCCATAGCGA 252 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAATGCA CCAACTGAA GTTTCTTTGG 256 GTTTTTCAAG ACTTATATC CCAAAGAATA CCGTTATATC ATACAACAAG CTTGTTTTG 256 GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTTATGAT AATTGACTACA 264 CTTATCCATG TCCCGGATTG CAAATGTTT TGATAATCAA CAACCATACT TATCAATCAA AATTAAACAA 2700 CCTGACTCCT CTAGGAATTCA GGAATCAGT TACCATCATA TATCTTTTTT TATTGACTGT 276 CTTATCCATG GGAGCCTT TACCACCTTAA GAATACATC TATCTTTTAT TATTGACTGT 276 CTTATTCCAG GGAGCCTT TATCTTCAT TAACCGTTCTA AACTTGCTAA AATTGCTCAA CACCTTCAA AACTTGC							
CAGGGTGTTC AAGGCTACCT TGATGGAAAA GGCTCTCTTA GAGATATTTG TAAATGGTAT 144 GATATCTCAA GTCGCTCTGT TCTCCAAAAG TGGATAAAAC GGTATACTAG TGGTGAAGAC 150 TTGAAAGCCA CTAGTAGAGG ATATAGCCGT ATGAAACAG GAAGGCAAGC CACATTTGAA 156 GAACGTGTAG AGATTGTTAA CTACACCATT GCCCATGGGA AAGACTATCA AGCAGCTATT 162 GAACGTGTAG AGATTGTTAA CTACACCATT TATTCTTGGG TGCGTAAACT TGAATGAAGAA 168 GGCTCACAAG GTTTGGTTG TAGACGTGTG AAAGGGTTAG AGAGTAACC TGATTTAACC 174 GAGATTGAC AACTTTAACT CAAGATTAAA CAATTGGAG AACGTAATCG TCTCTTAGAA 180 ATCGAGGTTA GTTTACTAAA AAAGTTAGAA GACATCAAAC GAGGAAACAG ACGGTAAGAC 186 GAGATGAGC TTTACCAAA AAAGTTAGAA GACATCAAAC GAGGAAACAG ACGGTAAGAC 186 TAGGTAAGCA TTTACCAAGAA TCAAAGAATTA TTACGATGAG GAATCTAATG 192 TGCCTATTCA GGCCTTATGC CAACTCTTGA AGGGGTCCG TCAGGCTAT TACAAGTGGC 198 TCAATCGTCA AAAAACAGAT TTTGAGACAA AAAATACAAA GCTAATGGCT AAAAACAGAA TTTATAAACA AACTTCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAACC 204 AACTTCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAACC 222 AAGAAAATAT TCTTAATCGT GAATTTACAG CCACAGCTCA TAACCAGAAA TGGTGCACAG 228 ATGTCAACCTA TCTTCAATAC GGTTAGCAAG CTATGACAACA ATTCTGGGGA 246 CTAAACCATA TCTTCAATAC GGTTAGCAAG CTAAAGCTTA TCTCAAGGAA TGGTGCACAG 228 ATGTCAACCTA TCTTCAATAC GGTCTGGGAG CTAAAGCTTA TCTCAGTGCG ATTAAAGACC 234 CTAATAACCG TCTCATATAC GCTTATGAGA TCCAGGAGCC ACACCTATCA TCCATAGCGA 246 CTAAGCCATT AAAAAGGGGC TAGAGCTCAA TCCAGGGAGC ACACCTATCA TCCATAGCGA 246 CTTATCCATG TCCCGGATTG GCAAAGAATA CCGTTATATC ATACAACAAG CTGGTCGAC 250 CTTATCCATG TCCCGGATTG GCAAATGTAT TCAAGGAGC ACACCTATCA TCCATAGCGA 246 CTTATCCATG TCCCGGATTG AAATCTAT TCAAGAGAAC TTTTTTGGCACA CAACCATACA AATTAAACAA 270 CCTGACTCCT CTAGAATCTA AACCACCTTAA GAAATACAC TCTTATGATA AATTAAACAA 270 CCTGACTCCT CTAGAATTCA GAAATCAGT TACAACCAC TATCATTCTAA AATTAAACAA 270 CCTGACTCCT CTAGAATTCA GAAATCAGT TACAACCAC TATCATACAA AATTAAACAA 270 CCTGACTCCT CTAGAATTCA GAAATCAGT TACAACCAC TACCTCTAA AATTAAACAA 270 CCTGACTCCT CTAGAATTCA GAATCAGT TACAACCAT AATTTGCTAA AATTAAACAA 270 CCTGACTCCT CTAGAATTCA GAATCAGT TACAACTAA TTCATCAGTT 276 CTACTTGACA GGGACCGTT CAGATCAGT TACACTCTA AATTTG	TGGAAGTAAT	ATTGCAAAGT	AATATATTTC	CTATTATATG	TTTATACGAT	ATTCGATATT	1320
TIGARAGCCA CTAGTAGAGG ATATAGCCGT ATGAAACAC GAACTATCA AGCACTATTGAA TIGAAAGCCA CTAGTAGAGG ATATAGCCGT ATGAAACAAG GAAGGCAAGC CACATTTGAA GAACGTGTAG AGATTGTTAA CTACACCATT GCCCATGGGA AAGACTATCA AGCAGCTATT GAGAAGGTTTG GTGTTCCTA CCAACAAATT TATTCTTGGG TGCGTAAGCT TGAGAAGAAA GAGATGTTG GTGTTCCTA CCAACAAATT TATTCTTGGG TGCGTAAGCT TGAGAAGAAA GGCTCACAAG GTTTGGTTGA TAGACCGTTG AAAGGGTTGG AGAGTAGGCC TGATTTAACC GAGAATTGAGC ACTTTAACT CAAGATTAAA CAATTGGAGG AACGTAATCG TCTCTTAGAA ATCGAGGTTA GTTTACTAA AAAGTTAGAA GACATCAAAC GAGGAAACAG ACGGTAAGAC TAGGTAAGCA TTTAGCGGAG TTCCAAGTAA TAAGGAATTA TTACGATGAG GAATCTAATG TGCCTATTCA GGCCTTATCC CAACTCTTGA AGGGGTCCG TTCAGGCTAT TACAAGTGGC TCAATCGTCA AAAAACAGAT TTTGAGACAA AAAATACAAA GCTAATGGCT AAAATCAAGG AACTTCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAATC GTCAACTTGG GACAACTTAA AACAAGAAAC GGATTCGTG ATTGATGACA ATTCTGGGGA AACTTCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAATC GTCAACTTGG GACAACTTAA AACAAGAAAC GGATTCGTG ATTGATGAAC ATTCTGGGGA AAGAAAAATAT TCTTAATCGT GTTAGCCATG CTTGTACAAA AGCTGGTGA AGATTTTACG CTAGTTCACCTA TCTTCAATAC GGTCTGGGGG CTAAAGCTTA TCCAAGGAAA TGGTGCACAG AAGAAAAATAT TCTTAATCG GAATTTACAG CCACAGCTCA TAACCAGAAA TGGTGCACAG CTATATCACCTA TCTTCAATAC GGTCTGGGGG CTAAAGCTTA TCCAAGTACA CTCATTGAGAA AGGAACCATT AAAAAGGGGC TAGAGCTCAA TCCAAGGACC ACCCTATCA TCCATAGCGA ATGGTCACCTA TCTTCAATAC GCTTATGAGA TCCAGGAGCC ACACCTATCA TCCATAGCGA CTTATCCACT TCCAGGATCT ACACCTTAA GAAATACAAC CCAACCTGAAC GTTCTTTTGG GAAGACCATT AAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCTATCA TCCATAGCGA CTTATCCACT TCCAGGATCT ACACCTTAA GAAATACAAC TCTTATGATG AGTTGGTCAA CTTATCCACT TCCAGGATCT ACACCTTAA GAAATACAAC TCTTATGATG AGTTGGTCAA CTTATCCACT TCCAGGATCT ACACCTTAA GAAATACAAC TCTTTATGAT AGTTCTTTGG GTTTTTCAAG ACTGAGATTCA ACACCTTAA GAAATACAAC TCTTTATTAT ATTTAACCTGT TGATGTGGCA CGTTATATCG AATTCTACAA CACACAACGT TATCAATCAA AATTAAACAA CCTGACTCCT CTAGAATTCG AGTTCTTACAA CACACAACGT TATCAATCAA AATTAAACAA CCTGACTCCT CTAGAATTCG AGTTCTTA ACCCTTTAA AATTTGCTAA AATGCTACA CAAAAATTAG CGGGGCCTT TAGTTTCTT TAACCGATCA	CCCACCCGTT	GTCGCGTTTA	CGGAAATACG	CCATTGATAT	ACTCCACATT	AGCTAAAGAA	1380
TTGAAAGCCA CTAGTAGAGG ATATAGCCGT ATGAAACAAG GAAGGCAAGC CACATTTGAA 156 GAACGTGTAG AGATTGTTAA CTACACCATT GCCCATGGGA AAGACTATCA AGCAGCTATT 162 GAGAAGTTTG GTGTTTCCTA CCAACAAATT TATTCTTGGG TGCGTAAGCT TGAGAAGAAT 168 GGCTCACAAG GTTTGGTTGA TAGACGTGTG AAAGGGTTGG AGAGTAGGCC TGATTTAACC 174 GAGATTGAGC AACTTTAACT CAAGATTAAA CAATTGGAGG AACGTAATCG TCTCTTAGAA 180 ATCGAGGTTA GTTTACTAAA AAAGTTAGAA GACATCAAAC GAGGAAACAG ACGGTAAGAC 186 TAGGTAAGCA TTTAGCGGAG TTCCAAGTAA TCAAGAATTA TTACGATGAG GAATCTAATG 192 TGCCTATTCA GGCCTTATGC CAACTCTTGA AGGGGTCCG TTCAGGCTAT TACAAGTGGC 198 TCAATCGTCA AAAAACAGAT TTTGAGACAA AAAATACAAA GCTAATGGCT AAAATCAAGG 204 AACTTCGTAG ACTCTACAAT GGTATCTTGA GTTATCGCCG TATGACAACA TTTATTAATC 210 GTCAACTTG GACAACTTAA AACAAGAAAC GTTATCGCGG TATGACAACA TTTATTAACC 220 GTCAACTTG GACAACTTAA AACAAGAAAC GTTGATGAAA AGCTGGTGA ATTCTGGGGA 216 GTAAACTGC CATTCGTCGT GTTAGCCATG CTTGTACAAA AGCTGGTGA ATTCTGGGGA 216 GTAAACTG CATTCGTCGT GTTAGCCATG CTTGTACAAA AGCTGGTGA ATTCTGGGGA 220 AAGAAAAAATA TCTTAATCGT GAATTTACAG CCACAGCTCA TAACCAGAAA TGTTAAGAGC 220 ATGTAAACGG TTCTAATACC GGTCTGGGAG CTAAAGCTTA TCTCAGAGAAC CATTGTTAT 240 GAAGACCATT AAAAAGGGC TAGAGCTCAA TCCAGGCCC ACACCTATCA TCCATAGCGA 246 GAAGACCATT AAAAAGGGGC TAGAGCTCAA TCCAGGGCC ACACCTATCA TCCATAGCGA 246 CTTATCCATG TCCCGGATTG GCAAAGAATA CCGTTATATC ATACAACAAG CTGGTCTGAC 250 CTTATCCATG TCCCGGATTG GCAAAGATA CCGTTATACA CACCTATCA TCCATAGCGA 264 CTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTCTTTGG 250 GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATA AATTAACACA 270 CCTGACTCCT CTAGAATTCA GAAATGCAT TACAACAAC TCTTATGATA AATTAACACA 270 CCTGACTCCT CTAGAATTCA GAAATGCAT TACAACAAC TCTTATGATA AATTAACACA 270 CCTGACTCCT CTAGAATTCA GAATTCTACAA CACACAACGT TATCAATCAA AATTAACACA 270 CCTGACTCCT CTAGAATTCA GAATTCTTA TACCTGTCT ACCTTCATACA AATTAACTA AATTAACACA 270 CCTGACTCCT CTAGAATTCA GAATTCTTA TACCTGTCT AACTTTTATT ATTTGACTAA AATAGCTACA 280 AAAAATTGAG CGAATCAAA AGCTTTCAT TACCTGTCT TCCCTCGACC 280 CAAAAAATGAG CCATTAAAA AACCTGGCT ATTTTTCCT CCTAAAAATT ATCTTCATA TACAACACAC TCTCCTCGACC 280 CAAAAAAA	CAGGGTGTTC	AAGGCTACCT	TGATGGAAAA	. GGCTCTCTTA	GAGATATTTG	TAAATGGTAT	1440
GAACGTGTAG AGATTGTTAA CTACACCATT GCCCATGGGA AAGACTATCA AGCAGCTATT 162 GAGAAGTTTG GTGTTTCCTA CCAACAAATT TATTCTTGGG TGCGTAAGCT TGAGAAGAAAT 168 GGCTCACAAG GTTTGGTTGA TAGACCGTGT AAAGGGTTGG AGAGTAAGCC TGATTTAACC 174 GAGATTGAGC AACTTTAACT CAAGATTAAA CAATTGGAGG AACGTAATCG TCTCTTAGAA 180 ATCGAGGTTA GTTTACTAAA AAAGTTAGAA GACATCAAAC GAGGAAACAG ACGGTAAGAC 186 GTGCCTATTCA GGCCTTATGC CAACTCTTGA AGGGGTCCG TTCAGGATGA CAACTCATCA GGCCTATTCA GGCCTATTCA GGCCTATCA GGCCTATTCA GGCCTTATGC CAACTCTTGA AGGGGTCCG TTCAGGCTAT TACCAAGTGGC 198 CTCAATCGTCA AAAAACAGAT TTTGGAGACAA AAAATACAAA GCTAATGGCT AAAATCAAGG 204 AACTTCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAATC 210 GTCAACTTGG GACAACTTAA AACAAGAAAC GGATTCGTTG ATTGAGAACA TTTTTTACGGCA 216 GTCAACTTGG GACAACTTAA AACAAGAAAC GGATTCGTTG ATTGAGAACA ATTCTGGGGA 216 GTCAACTTGG GACAACTTAA AACAAGAAAC GCATACAAAA AGCTGGTGAC AGATTTTACG 222 AAGAAAAAATAT TCTTAATCGT GAATTTACAG CCACAGCTCA TAACCAGAAA TGGTGCACAG 228 ATGTCACCTA TCTCAATAC GGTCTGGGAG CTAAAGCTTA TCCCAGAAA TGGTGCACAG 228 ATGTCACCTA TCTTCAATAC GGTCTGGGAG CTAAAGCTTA TCCCAGAAA TGGTGCACAG 228 ATGTAAACGG TTCTATATC GCTTATGAGA TAGCCACAA CAATGAAATC CACTTGTTAT 240 GAAGACCATT AAAAAGGGC TAGAGCTCAA TCCAGGGCC ACACCTATCA TCCATAGCGA 246 CTTATCCATG TCCCGGATTG GCAAAGATA CCGTTATATC ATACCAGAAA GTTTCTTGG GTTTTCCAG ACTGGAGTCT ACCACCTTAA GAAATACAAC CCAACTGAAA GTTTCTTTGG 256 GTTTTTCAAG ACTGGATTC ACCACCTTAA GAAATACAAC TCTTATGATA AGTTGCTCAC 250 CTTATCCATG TCCCGGATTG ACACCTTAA GAAATACAAC TCTTATGATA AATTAACAA 270 CCTGACTCCT CTAGAATTCA GAATTCTACAA CACACAACGT TATCAATCAA AATTAACACA 270 CCTGACTCCT CTAGAATTCA GAATTCTACAA CACACAACGT TATCAATCAA AATTAACACA 270 CCTGACTCCT CTAGAATTCA GAATTCTTCT TAACCTGTT ACCTTTTTTT ATTTTTTTTTT	GATATCTCAA	GTCGCTCTGT	TCTCCAAAAG	TGGATAAAAC	GGTATACTAG	TGGTGAAGAC	1500
GAGAAGTTTG GTGTTTCCTA CCAACAAATT TATCTTGGG TGCGTAAGCT TGAGAAGAAT 168 GGCTCACAAG GTTTGGTTGA TAGACGTGTG AAAGGGTTGG AGAGTAAGCC TGATTTAACC 174 GAGATTGAGC AACTTTAACT CAAGATTAAA CAATTGGAGG AACGTAATCG TCTCTTAGAA 180 ATCGAGGTTA GTTTACTAAA AAAGTTAGAA GACATCAAAC GAGGAAACAG ACGGTAAGAC 186 TAGGTAAGCA TTTAGCGGAG TTCCAAGTAA TCAAGAATTA TTACGATGAG GAATCTAATG 192 TGCCTATTCA GGCCTTATGC CAACTCTTGA AGGGGTCTCG TTCAGGCTAT TACAAGTGGC 198 TCAATCGTCA AAAAACAGAT TTTGAGACAA AAAATCAAAA GCTAATGGCT AAAATCAAGG 204 AACTCCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAATC 210 GTCAACTTGG GACAACTTA AACAAGAAAC GGATTCGTTG ATTGATGACA ATTCTGGGGA 216 TTAGTTCAGT CATTCGTCGT GTTAGCCATG CTTGTACAAA AGCTGGTGAC AGATTTTACG 222 AAGAAAAATT TCTTAATCGT GAATTTACAG CCACAGCTCA TAACCAGAAA TGGTGCACAG 228 ATGTCACCTA TCTTCAATAC GGTCTGGGAG CTAAAGCTTA TCTCAGTGCG ATTAAAGACC 234 CGTATAAACGG TTCTATTATC GCTTATGAGA TTAGTCACAA CAATGAAATC CACTTGTTAT 2400 GAAGACCATT AAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCCTATCA TCCATAGCGA 2460 CTTATCCATG TCCCGGATTG GCAAATGTAT TCGAGGAGC ACACCTATCA TCCATAGCGA 2600 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAAACAA CATTCTTAGA GTTTCTTTGG CTTTTTCAGA ACTGAGACCATT ACCACGATTA ACCACCTATA ACAACAAGA TTCTTTTGG CTTTTTCAGA ACTGAGACCATT ACCACGATA TCCAGGAGCC ACACCCTATCA TCCATAGCGA 2600 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAAACAA CAATGAAATC CACTTGTTAT 2600 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAAACAA CTTTATGATCGA ACTTCTTTGG CTTTTTCAGA ACTGAGTCAA ACCACCTATCA ACTTCTTTAGA ACTTGGTCAA ACTTCATAACAA CACACAACGT TATCAATCAA AATTAAACAA 2700 CCTGACTCCT CTAGAATTCA GAAATCAGT TATCATCAA AATTAAACAA 2700 CCTGACTCCT CTAGAATTCA GAAATCAGT TATCTTTTAT ATTTGACTGT 2760 CTACTTGACA GGGGCCTT CAGAATCAGT TGCATAACTT ATCTTTTAT ATTTGACTGT 2760 CACTTGACCAC CCACTTAAACT TACCTTTATATC AATTTCAATCAA	TTGAAAGCCA	CTAGTAGAGG	ATATAGCCGT	ATGAAACAAG	GAAGGCAAGC	CACATTTGAA	1560
GGCTCACAAG GTTTGGTTGA TAGACGTGTG AAAGGGTTGG AGAGTAGGCC TGATTTAACC 174 GAGATTGAGC AACTTTAACT CAAGATTAAA CAATTGGAGG AACGTAATCG TCTCTTAGAA 18.0 ATCGAGGGTTA GTTTACTAAA AAAGTTAGAA GACATCAAAC GAGGAAACAG ACGGTAAGAC 18.6 TAGGTAAGCA TTTAGCGGAG TTCCAAGTAA TCAAGAATTA TTACGATGAG GAATCTAATG 19.2 TGCCTATTCA GGCCTTATGC CAACTCTTGA AGGGGTCTCG TTCAGGCTAT TACAAGTGGC 19.8 TCAATCGTCA AAAAACAGAT TTTGAGACAA AAAATCAAAA GCTAATGGCT AAAATCAAGG 20.4 AACTTCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAATC 21.0 GTCAACTTGG GACAACTTAA AACAAGAAAC GGATTCGTTG ATTGATGACA ATTCTGGGGA 21.6 TTAGTTCAGT CATTCGTCGT GTTAGCCATG CTTGTACAAA AGCTGGTGAC AGATTTACG 22.2 AAGAAAATAT TCTTAATCGT GAATTTACAG CCACAGGCTCA TAACCAGAAA TGGTGCACAG 22.8 ATGTCACCTA TCTTCAATAC GGTCTGGGGA CTAAAGCTTA TCTCAGTGCG ATTAAAGACC 23.4 TGTATAACGG TTCTATTATC GCTTATGAGA TTAGTCACAA CAATGAAAAT CACTTGTTAT 24.0 GAAGACCATT AAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCTATCA TCCATAGCGA 25.2 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATGACA CAATGAAAAC CTGGTCTGAC 25.2 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATCACAA CAATGAAAAC CTGGTCTGAC 25.2 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATCACAA CAATGAAAAC CTGGTCTGAC 25.2 CTTATCCATG TCCCGGATTG ACACCCTTAA GAAATACAA CTCTTATGATG AGTTCTTTGG 25.2 CTTATCCATG TCCCGGATTG AAATCTTA ACACCACAACGT TATCAATCAA AATTAAACAA 27.0 CCTGACTCCT CTAGAATTCA GAAATCAGT TATCATACAA AATTAAACAA 27.0 CCTGACTCCT CTAGAATTCA GAAATCAGT TACAACAACGT TATCAATCAA AATTAAACAA 27.0 CCTGACTCCT CTAGAATTCA GGAATCAGGT TGCATAACTT ATCTTTTATT ATTTGACTGT 27.6 CTACTTGACA GGGAGCCTT CAGATTCCTT AACCTTCTA AATTTGCTAA AATTAACCAA 28.2 AAAAAATGAG CCATTTAATG CTTATTTCTT AACCTTCTA AATTTGCTAA AATTAACCAA 28.2 AAAAAATGAG CCATTTAATG CTTATTTCTT AACCTTCTA AATTTCCATA TTCATCAGTT 29.4 AAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 29.4 AAAAATTGAG CGTGAGGCTT TTTGTTTCAT AACCTTCTA AATTTCCATA TTCATCAGTT 29.4 AAAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 29.4 AAAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 29.4 AAAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 29.4 AAAAA	GAACGTGTAG	AGATTGTTAA	CTACACCATT	GCCCATGGGA	AAGACTATCA	AGCAGCTATT	1620
ARGATTGAGC AACTTTAACT CAAGATTAAA CAATTGGAGG AACGTAATCG TCTCTTAGAA 1800 ATCGAGGTTA GTTTACTAAA AAAGTTAGAA GACATCAAAC GAGGAAACAG ACGGTAAGAC 1860 TAGGTAAGCA TTTAGCGGAG TCCCAAGTAA TCAAGAATTA TTACGATGAG GAATCTAATG 1920 TGCCTATTCA GGCCTTATGC CAACTCTTGA AGGGGTCTCG TTCAGGCTAT TACAAGTGGC 1980 TCAATCGTCA AAAAACAGAT TTTGAGACAA AAAATACAAA GCTAATGGCT AAAATCAAGG 2040 AACTTCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAATC 2100 GTCAACTTGG GACAACTTAA AACAAGAAAC GGATTCGTTG ATTGATGACA ATTCTGGGGA 2160 TTAGTTCAGT CATTCGTCG GTTAGCCATG CTTGTACAAA AGCTGGTGAC AGATTTACG 2220 AAGAAAAATAT TCTTAATCGT GAATTTACAG CCACAGCTCA TAACCAGAAA TGGTGCACAG 2280 ATGTCACCTA TCTTCAATAC GGTCTGGGGA CTAAAGCTTA TCTCAGTGCG ATTAAAGACC 2340 TGTATAACGG TTCTATTATC GCTTATGAGA TTAGTCACAA CAATGAAATC CACTTGTTAT 2400 GAAGACCATT AAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCTATCA TCCATAGCGA 2460 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAAACAC CCACACTACA TCCATAGCGA 2500 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAACAC CCACACTACA TCCATAGCGA 2640 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAAACAC CCACACTGAAA GTTTCTTTGG 2580 GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAC TCTTATGATG AGTTGGTCAA 2640 TGATGTGGCA CGTTATATCG AATTCACAA CACACCAACGT TATCAATCAA AATTAAACAA 2700 CCTGACTCCT CTAGAATTCA GGAATCAGGT TGCATAACCTT ATCTTATTAT ATTTGACTGT 2760 CTACTTGACA GGGAGCCGTT CAGATTCGTT AACCCTTCA AATTTGCTAA AATAACACA 2820 AAAAAATGAG CCATTTAATG CTTATTCTT ATACTGTTTTA ATTTGACTGT 2760 CTACTTGACA GGGAGCCGTT CAGATTCGTT AACCTTTCTA AATTTGCTAA AATAACCAA 2820 AAAAAATAGA CGGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 2940 AAAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 2940 CTAAAAAATAA AACCTGAAAAAAAAAAAAAAAAAAAAAA	GAGAAGTTTG	GTGTTTCCTA	CCAACAAATT	TATTCTTGGG	TGCGTAAGCT	TGAGAAGAAT	1680
ATCGAGGTTA GTTTACTAAA AAAGTTAGAA GACATCAAAC GAGGAAACAG ACGGTAAGAC 1866 TAGGTAAGCA TTTAGCGGAG TTCCAAGTAA TCAAGAATTA TTACGATGAG GAATCTAATG 1926 TGCCTATTCA GGCCTTATGC CAACTCTTGA AGGGGTCTCG TTCAGGCTAT TACAAGTGGC 1986 TCAATCGTCA AAAAACAGAT TTTGAGACAA AAAATCAAA GCTAATGGCT AAAATCAAGG 2046 AACTTCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAATC 2106 GTCAACTTGG GACAACTTAA AACAAGAAAC GGATTCGTTG ATTGATGAAC ATTCTGGGGA 2166 TTAGTTCAGT CATTCGTCGT GTTAGCCATG CTTGTACAAA AGCTGGTGAC AGATTTACG 2226 AAGAAAATAT TCTTAATCGT GAATTACAG CCACAGCTCA TAACCAGAAA TGGTGCACAG 2286 ATGTCACCTA TCTTCAATAC GGTCTGGGAG CTAAAGCTTA TCTCAGTGCG ATTAAAGACC 2340 TGTATAACGG TTCTATTATC GCTTATGAGA TCAGAGACC ACACCTATCA TCCATAGCGA 2460 GAAGACCATT AAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCTATCA TCCATAGCGA 2460 TTGAGGTAGT CAATATACTT CCAAAGAATA CCGTTATATC ATACAACAAG CTTGTTTATC GCTTATCCATG TCCCGGATTG GCAAATGTAT TGATAATCAC CCAACCTGAAA GTTTCTTTGG 2560 GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATCAAAC TCTTATGATA AATTAAACAA 2700 CCTGACTCCT CTAGAATTCA GGAATCAGT TGCATAACCT TATCTATGAT ACTTTTTTGG 2560 CCTGACTCCT CTAGAATTCA GGAATCACT TACAACAACGT TATCAATCAA AATTAAACAA 2700 CCTGACTCCT CTAGAATTCA GGAATCAGT TGCATAACCT TATCTTTATT ATTTGACTGT 2760 CTACTTGACA GGGAGCCGTT CAGATTCATT AACCTTCTA AATTTGCTAA AATAGCTACA 2820 AAAAAATGAA GCGTGAGGCTT TTTGTTTCAT TAACCGATCA TCCTCCGGACC 2880 AAAAAATTGAG CCATTAAATG CTTATTTCTT TAACCGTTTT GCCTCACGCT CTCCTGGACC 2880 AAAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAACCGATTA ATTTTCCATA TTCATCAGTT 2940 TGTTTTCCGA GAGCCATCAA AGCTTCGATA AGGTTGATA TTCCAGGAAT AAAGGTAATA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGGCGCTCCA 3000	GGCTCACAAG	GTTTGGTTGA	TAGACGTGTG	AAAGGGTTGG	AGAGTAGGCC	TGATTTAACC	1740
TAGGTAAGCA TTTAGCGGAG TTCCAAGTAA TCAAGAATTA TTACGATGAG GAATCTAATG 1920 TGCCTATTCA GGCCTTATGC CAACTCTTGA AGGGGTCTCG TTCAGGCTAT TACAAGTGGC 1980 TCAATCGTCA AAAAACAGAT TTTGAGACAA AAAATCAAAA GCTAATGGCT AAAATCAAGG 2040 AACTTCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAATC 2100 GTCAACTTGG GACAACTTAA AACAAGAAC GGATTCGTTG ATTGATGACACA ATTCTGGGGA 2160 GTCAACTTGG GACAACTTAA AACAAGAAAC GGATTCGTTG ATTGATGACACA ATTCTGGGGA 2160 GTCAACTTGG GACAACTTAA AACAAGAAAC GGATTCGTTG ATTGATGACAAC ATTCTGGGGA 2220 AAGAAAAATAT TCTTAATCGT GAATTTACAG CCACAGGCTCA TAACCAGAAA TGGTGCACAG 2280 ATGTCACCTA TCTTCAATAC GGTCTGGGAG CTAAAGCTTA TCTCAGTGCG ATTAAAGACC 2340 GGAAGACCATT AAAAAAGGGGC TAGAGCTCAA TCCAGGAGC ACACCTATCA TCCATAGCGA 2460 GAAGACCATT AAAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCTATCA TCCATAGCGA 2460 GTTATCCATG TCCCGGATTG GCAAATGTAT TGATAATGCA CCAACTGAAA GTTTCTTTGG 2580 GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTCTTTGG 2580 GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTGGTCAA 2640 TGATGTGGCA CGTTATATCG AATTCTACAA CACCACAACGT TATCAATCAA AATTAAACAA 2700 CCTGACTCCT CTAGAATTCA GGAATCAGT TGCATAACCT TCTTATGATG AGTTGGTCAA 2640 CCTGACTCCT CTAGAATTCA GGAATCAGT TGCATAACTT ATCTTTTATT ATTTGACTGT 2760 CTACTTGACA GGGGCCGTT CAGATTGCTT AACCTTCTTA AATTTTGCTAA AATAGCTACA 2820 AAAAAATGAG CCATTTAATG CTTATTTCTT AACCTTTCTA AATTTTCCATA TTCATCAGCT 2880 AAAAAATGAG CCATTTAATG CTTATTTCTT TAAACGATGA TATTTCCATA TTCATCAGTT 2940 TGTTTTCCAG GAGCCGTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 2940 TGTTTTCCAG GAGCCATCAA AGCTTCGATA AGGTTGATA AAGGTAATA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGGCGCTCCA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTTCCTG CGTAAAAATTT ATGGCGCTCCA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTTCCTG CGTAAAAATTT ATGGCGCTCCA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTTCCTG CGTAAAAATTT ATGGCGCTCCA 3000 CTAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA	GAGATTGAGC	AACTTTAACT	CAAGATTAAA	CAATTGGAGG	AACGTAATCG	TCTCTTAGAA	1800
TECCTATTCA GGCCTTATGC CAACTCTTGA AGGGGTCTG TTCAGGCTAT TACAAGTGGC 1980 TCAATCGTCA AAAAACAGAT TTTGAGACAA AAAATACAAA GCTAATGGCT AAAATCAAGG 2040 AACTTCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAATC 2100 GTCAACTTGG GACAACTTAA AACAAGAAAC GGATTCGTTG ATTGATGAAC ATTCTGGGGA 2160 TTAGTTCAGT CATTCGTCGT GTTAGCCATG CTTGTACAAA AGCTGGTGAC AGATTTTACG 2220 AAGAAAATAT TCTTAATCGT GAATTTACAG CCACAGCTCA TAACCAGAAA TGGTGCACAG 2280 ATGTCACCTA TCTTCAATAC GGTCTGGGAG CTAAAGCTTA TCTCAGTGCG ATTAAAGACC 2340 GGAAGACCATT AAAAAGGGGC TAGAGCTCAA TCCAGGAAC CACTCATCA TCCATAGCGA 2460 TTGAGGTAGT CAATATACTT CCAAAGAATA CGGTTATATC ATACAACAAG CTGGTCTGAC 2520 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAATGCA CCAACTGAAA GTTTCTTTGG 2580 GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTCTTTGG 2580 GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTGGTCAA 2640 CCTGACTCCT CTAGAATTCA GAATCACAA CACCACACCT TATCCATGAA AATTAAACAA 2700 CCTGACTCCT CTAGAATTCA GAATCACAA CACCACACCT TATCCATCA AATTAAACAA 2700 CCTGACTCCT CTAGAATTCA GAATCACAT TACATCAAA AATTAAACAA 2700 CCTGACTCCT CTAGAATTCA GAATCACGT TGCATAACCTT ATCTTTTATT ATTTGACTGT 2760 CTACTTGACA GGGAGCCGTT CAGATTGCTT AACCTTTCTA AATTTGCTAA AATTAACCAA 2800 CCTGACTCCT CTAGAATTCA GAATCACTT AACCTTTCTA AATTTGCTAA AATTAACCAA 2800 AAAAATTGAG CCATTTAATG CTTATTTCTT ATACTGTCTT GCCTCACGCT CTCCTCGACC 2880 AAAAAATGAG CCATTTAATG CTTATTTCTT ATACTGTCTT GCCTCACGCT CTCCTCGACC 2800 AAAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 2940 TGTTTTCCGA GAGCCATCAA AGCTTCGATA AGGTCGATAA TTCCAGGAAT AAAGGTAATA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTCCTG CGTAAAAATTT ATGCGCTCCA 3000	ATCGAGGTTA	GTTTACTAAA	AAAGTTAGAA	GACATCAAAC	GAGGAAACAG	ACGGTAAGAC	1860
TCAATCGTCA AAAAACAGAT TTTGAGACAA AAAATACAAA GCTAATGGCT AAAATCAAGG 2040 AACTTCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAATC 2100 GTCAACTTGG GACAACTTAA AACAAGAAAC GGATTCGTTG ATTGATGACA ATTCTGGGGA 2160 TTAGTTCAGT CATTCGTCGT GTTAGCCATG CTTGTACAAA AGCTGGTGAC AGATTTTACG 2220 AAGAAAATAT TCTTAATCGT GAATTTACAG CCACAGCTCA TAACCAGAAA TGGTGCACAG 2280 ATGTCACCTA TCTTCAATAC GGTCTGGGAG CTAAAGCTTA TCTCAGTGCG ATTAAAGACC 2340 TGTATAACGG TTCTATTATC GCTTATGAGA TCAGAGACC ACACCTATCA TCCATAGCGA 2460 GAAGACCATT AAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCTATCA TCCATAGCGA 2520 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAATGCA CTAACACAGA GTTTCTTTGG 2580 GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAC TCTTATGATG AGTTGGTCAA 2640 TGAGTGGGCA CGTTATATCG AATTCTACAA CACACAACGT TATCAATCAA AATTAAACAA 2700 CCTGACTCCT CTAGAATTCA GGAATCAGGT TGCATAACTT ATCTTTTATT ATTTGACTGT 2760 CTACTTGACA GGGAGCCGTT CAGATTGCTT AACCTTCTA AATTTGCTAA AATAACCAA 2820 AAAAAATGAG CCATTTAATG CTTATTTCTT AACCTTTCTT GCCTCCAGCC 2880 AAAAAATGAG CCATTTAATG CTTATTTCTT AACCTTCTT AATTTCCATA TTCATCAGTT 2940 TGTTTTCCGA GAGCCGTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 2940 TGTTTTCCGA GAGCCATCAA AGCTTCGATA AGGTCGATA TTCCAGGAAT AAAGGTAATA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCTCCA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCTCCA 3000 CTAAAAAATAAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCTCCA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCTCCA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCTCCA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCTCCA 3000 CTAAAAAATAAA TATATAAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCCTCCA 3000 CTAAAAAATAAA TATATAAAAAA AACCTGGCCT ATTTTTCCTG CGT	TAGGTAAGCA	TTTAGCGGAG	TTCCAAGTAA	TCAAGAATTA	TTACGATGAG	GAATCTAATG	1920
AACTTCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAATC 2100 GTCAACTTGG GACAACTTAA AACAAGAAAC GGATTCGTTG ATTGATGACA ATTCTGGGGA 2160 TTAGTTCAGT CATTCGTCGT GTTAGCCATG CTTGTACAAA AGCTGGTGAC AGATTTTACG 2220 AAGAAAATAT TCTTAATCGT GAATTTACAG CCACAGCTCA TAACCAGAAA TGGTGCACAG 2280 ATGTCACCTA TCTTCAATAC GGTCTGGGAG CTAAAGCTTA TCTCAGTGCG ATTAAAGACC 2340 TGTATAACGG TTCTATTATC GCTTATGAGA TTAGTCACAA CAATGAAATC CACTTGTTAT 2400 GAAGACCATT AAAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCTATCA TCCATAGCGA 2460 CTTATCCATG CAATATACTT CCAAAGAATA CCGTTATATC ATACAACAAG CTGGTCTGAC 2520 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAATGCA CCAACTGAAA GTTTCTTTGG 2560 GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTGGTCAA 2640 CCTGACTCCT CTAGAATTCA GGAATCAGGT TGCATAACTT ATCTTTATT ATTTGACTGT 2760 CTACTTGACA GGGAGCCGTT CAGATTGCTT AACCTTTCTA AATTTGCTAA AATAGCTACA 2820 AAAAAATGAG CCATTTAATG CTTATTTCTT ATACTGTCTT GCCTCACGCT CTCCTCGACC 2880 AAAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TCCATCAGTT 2940 TGTTTTCCGA GAGCCATCAA AGCTTCGATA AGGTCGATAA TTCCACGATT AAAAGATAA AAAAGGTAATA AGCTTCCATA TAAACGAATGA TATTTCCATA TACAACAAT AAAAGGTAATA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTCCTG CGTAAAAATTT ATGCGCTCCA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT AAGCGCTCCA 30000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT AAGCGCTCCA 30000 CTAAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT AAGCGCTCCA 30000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT AAGCGCTCCA 30000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT AAGCGCTCCA 30000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT AAGCGCTCCA 30000 CTAAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT AAGCGCTCCA 300000000000000000000000000000000000	TGCCTATTCA	GGCCTTATGC	CAACTCTTGA	AGGGGTCTCG	TTCAGGCTAT	TACAAGTGGC	1980
GTCAACTTGG GACAACTTAA AACAAGAAC GGATTCGTTG ATTGATGAC ATTCTGGGGA 2220 TTAGTTCAGT CATTCGTCGT GTTAGCCATG CTTGTACAAA AGCTGGTGAC AGATTTTACG 2220 AAGAAAATAT TCTTAATCGT GAATTTACAG CCACAGCTCA TAACCAGAAA TGGTGCACAG 2280 ATGTCACCTA TCTTCAATAC GGTCTGGGAG CTAAAGCTTA TCTCAGTGCG ATTAAAGACC 2340 TGTATAACGG TTCTATTATC GCTTATGAGA TTAGTCACAA CAATGAAATC CACTTGTTAT 2400 GAAGACCATT AAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCTATCA TCCATAGCGA 2460 TTGAGGGTAGT CAATATACTT CCAAAGAATA CCGTTATATC ATACAACAAG CTGGTCTGAC 2520 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAATGCA CCAACTGAAA GTTTCTTTGG 2580 GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTGGTCAA 2640 TGATGTGGCA CGTTATATCG AATTCTACAA CACACAACGT TATCAATCAA AATTAAACAA 2700 CCTGACTCCT CTAGAATTCA GGAATCAGGT TGCATAACTT ATCTTTTATT ATTTGACTGT 2760 CTACTTGACA GGGAGCCGTT CAGATTGCTT AACCTTTCTA AATTTGCTAA AATTAGCTACA 2820 AGAAAAACGAG CCATTTAATG CTTATTTCTT ATACCTGTTT GCCTCACGCT CTCCTCGACC 2880 AAAAAATGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TACACGATT AAAGGTAATA 3000 TGTTTTCCGA GAGCCATCAA AGCTTCGATA AGGTCGATAA TTCCAGGAAT AAAGGTAATA 3000 CTAAAAAATAA TATAAAAAA ACCTGGCCT ATTTTCCTG CGTAAAAATTT ATGCGCTCCA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3000	TCAATCGTCA	AAAAACAGAT	TTTGAGACAA	AAAATACAAA	GCTAATGGCT	AAAATCAAGG	2040
TTAGTTCAGT CATTCGTCGT GTTAGCCATG CTTGTACAAA AGCTGGTGAC AGATTTACG 2220 AAGAAAATAT TCTTAATCGT GAATTTACAG CCACAGCTCA TAACCAGAAA TGGTGCACAG 2280 ATGTCACCTA TCTTCAATAC GGTCTGGGAG CTAAAGCTTA TCTCAGTGCG ATTAAAGACC 2340 TGTATAACGG TTCTATTACC GCTTATGAGA TTAGTCACAA CAATGAAATC CACTTGTTAT 2400 GAAGACCATT AAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCTATCA TCCATAGCGA 2460 CTTAGCGTAGT CAATATACTT CCAAAGAATA CCGTTATACC ATACAACAAG CTGGTCTGAC 2520 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAATGCA CCAACTGAAA GTTTCTTTGG 2580 GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTGGTCAA 2700 CCTGACTCCT CTAGAATTCA GGAATCAGGT TGCATAACTT ATCTTTATT ATTTGACTGT 2700 CTACTTGACA GGGAGCCGTT CAGATTGCTT AACCTTCTA AATTTGCTAA AATAGCTACA 2820 AGAAAAACGAG CCATTTAATG CTTATTTCTT ATACTGTTT GCCTCCACGCT CTCCTCGACC 2880 AAAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 2940 TGTTTTCCGA GAGCCATCAA AGCTTCGATA AGGTCGATAA TTCCAGGAAT AAAGGTAATA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3000 CTAAAAAATTAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3000 CTAAAAAATTA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3000 CTAAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCTCCA 3000 CTAAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCTCCA 3000 CTAAAAAATAA TATATAAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCTCAA 3000 CTAAAAATAAA TATTTAAAAAAAAAAAAAAAAAAAAA	AACTTCGTAG	ACTCTACAAT	GGTATCTTAG	GTTATCGCCG	TATGACAACA	TTTATTAATC	2100
AAGAAAATAT TCTTAATCGT GAATTTACAG CCACAGCTCA TAACCAGAAA TGGTGCACAG 228 CATGTCACCTA TCTTCAATAC GGTCTGGGAG CTAAAGCTTA TCTCAGTGCG ATTAAAGACC 234 CATGTATAACGG TTCTATTATC GCTTATGAGA TTAGTCACAA CAATGAAATC CACTTGTTAT 240 CAAGAGACCATT AAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCTATCA TCCATAGCGA 246 CATGTATCCATG TCCCGGATTG GCAAAGAATA CCGTTATATC ATACAACAAG CTGGTCTGAC 252 CATGTCTCATG TCCCGGATTG GCAAATGTAT TGATAATGCA CCAACTGAAA GTTTCTTTGG 258 CAACTGTCAC ACCCCTTATCAACAA ACTTAAACAA 270 CAACTGACA CACACTGAAA AATTAAACAA 270 CAACTGACA CACACTGACA AATTAAACAA 270 CAACTGACA GGGAGCCGT CAGAATTCA AACCATGCAT AACTTTTATT ATTTGACTGT 276 CAACAAACACG CAACTGACA CACACTGACA CACACTGACA AATTAAACAA 270 CAACATGACA GGGAGCCGT CAGAATTCAA AACCATTCAA AATTAAACAA AATAACCAA 282 CAACAAACGA CACACTGACA AATTAAACAA 270 CAACAACGA CACACTGACA AATTAAACAA 270 CAACATTGACA GGGAGCCGT CAGAATTCAA AACCATTCAA AATTTGACTGT 276 CAACAAACGA CACACAGA AATTAAACAA AATAACAA 270 CAACAAACGA CACACAGA AATTAAACAA AATAACAA 270 CAACAAACGA CAACGATAACTA AATTTGACTGA CAACAAACGA CAACAACGA AATTAAACAA AATAACAA 270 CAACAAACGA CAACAACGA AATTAAACAA AATAACAA 270 CAACAAACGA CAACAACGA AATTAAACAA AATAACAA 270 CAACAAACGA CAACAACGA AATTAAAAAA AACCTGCTT AAACCTTTCTA AATTTGCTAA AATAACAA AATAACAA 282 CAACAAAAAATTGAA CAACAAACGA CAACAAACGA CAACAACGT TATTTATTAAAAAA AACCTTGCTA AACCTTTCTA AATTTTCATAA AATAACAA 282 CAACAAAAAATTGAA CAACAAACGA CAACAAACGT TATTTTCATAAAAAAAATTAAAAAA AACCTGGACT AAAAAATTAAAAAA AACCTGGACT AAAAAATTAAAAAA AACCTGGACT ATTTTTCCTG CGTAAAAATTA AAAAGGTAAAAA AAAAAAAAAA	GTCAACTTGG	GACAACTTAA	AACAAGAAAC	GGATTCGTTG	ATTGATGAAC	ATTCTGGGGA	2160
ATGTCACCTA TCTTCAATAC GGTCTGGGAG CTAAAGCTTA TCTCAGTGCG ATTAAAGACC 234 CTGTATAACGG TTCTATTATC GCTTATGAGA TTAGTCACAA CAATGAAATC CACTTGTTAT 240 CTAAGACCAT AAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCTATCA TCCATAGCGA 246 CTTTTGAGGTAGT CAATATACTT CCAAAGAATA CCGTTATTATC ATACAACAAG CTGGTCTGAC 252 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAATGCA CCAACTGAAA GTTTCTTTGG 258 CTAATGTGAGAA ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTGGTCAA 264 CTGATGTGGAC CGTTATATCG AATTCTACAA CACACAACGT TATCAATCAA AATTAAACAA 270 CCTGACTCCT CTAGAATTCA GGAATCAGGT TGCATAACCT ATCTTTTATT ATTTGACTGT 276 CTACTTGACA GGGAGCCGTT CAGATTGCTT AACCTTTCTA AATTTGCTAA AATTAGACAA 282 CTACTTGACA GGGAGCCGTT CAGATTCCT AACCTTTCTA AATTTGCTAA AATAGCTACA 282 CTACTTGACA CGTGAGGCT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 294 CTACTTTCCGA GAGCCATCAA AGCTTCGATA AGGTCGATAA TTCCAGGAAT AAAGGTAATA 300 CCTAAAAATAA TATATAAAAAA AACCTGGCCT ATTTTCCTG CGTAAAATTT ATGCGCTCCA 30 CCTAAAAATAA TATATAAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 30 CCTAAAAAATAA TATATAAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 30 CCTAAAAATAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 30 CCTAAAAATAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 30 CCTAAAAATTT ATGCGCTCCA 30 CCTAAAATTT ATGCGCTCCAA 30 CCTAAAATTT ATGCGCTCCAA 30 CCTAAAATTT ATGCGCTCCAA 30 CCTAAAATTT ATGCGCTCCAA 30 CCTAAAATTT ATGCGCT	TTAGTTCAGT	CATTCGTCGT	GTTAGCCATG	CTTGTACAAA	AGCTGGTGAC	AGATTTTACG	2220
TGTATAACGG TTCTATTATC GCTTATGAGA TTAGTCACAA CAATGAAATC CACTTGTTAT 240 CGAAGACCATT AAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCTATCA TCCATAGCGA 246 CGAAGACACAT CAATATACTT CCAAAGAATA CCGTTATATC ATACAACAAG CTGGTCTGAC 252 CGTTATACCATG TCCCGGATTG GCAAATGTAT TGATAATGCA CCAACTGAAA GTTTCTTTGG 258 CGTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTGGTCAA 264 CGTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTGGTCAA 270 CCCGGACTCCT CTAGAATTCA GGAATCAGA TGCATAACCT ATCTTTATT ATTTGACTGT 276 CCTGACCTCCT CTAGAATTCA GGAATCAGGT TGCATAACCT ATCTTTTATT ATTTGACTGT 282 CAAAAAAACGAG CCATTTAATG CTTATTTCTT ATACTGTCTT GCCTCACGCT CTCCTGGACC 288 CAAAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 294 CCTGAAAAATAA AGCCTTCGATA AGCTTGCTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTCCTG CGTAAAAATT AAAGGATAATA 300 CCTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATT ATGCGCTCCA 300 CCTAAAAAATAA TATATAAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATT ATGCGCTCCA 300 CCTAAAAAATAA TATATAAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATT ATGCGCTCCA 300 CCTAAAAAATAA TATATAAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATT ATGCGCTCCA 300 CCTAAAAATTA ATGCGCTCCA 300 CCTAAAAATTA ATGCGCTCCA 300 CCTAAAAATTA ATGCGCTCCA 300 CCTAAAAATTA ATTTTTTCCTG CGTAAAAATTT ATGCGCTCCA 300 CCTAAAAATTA ATGCGAAAATTA ATGCGAAAATTA ATGCGCTCA ATTTTTTCCTG CGTAAAAATTA ATGCGCTCA 300 CCTAAAAATTA ATGCGAAAATTA ATGCGAAAATTA ATGCAAAATTA ATGCAAAATTA ATGCAAAATTA ATGCAAAAATTA ATGCAAAAATAAAAAAAAAA	AAGAAAATAT	TCTTAATCGT	GAATTTACAG	CCACAGCTCA	TAACCAGAAA	TGGTGCACAG	2280
GAAGACCATT AAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCTATCA TCCATAGCGA 24600 TTGAGGTAGT CAATATACTT CCAAAGAATA CCGTTATATC ATACAACAAG CTGGTCTGAC 25200 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAATGCA CCAACTGAAA GTTTCTTTGG 25800 GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTGGTCAA 26400 TGATGTGGCA CGTTATATCG AATTCTACAA CACACAACGT TATCAATCAA AATTAAACAA 27000 CCTGACTCCT CTAGAATTCA GGAATCAGGT TGCATAACTT ATCTTTATT ATTTGACTGT 27600 CTACTTGACA GGGAGCCGTT CAGATTGCTT AACCTTTCTA AATTTGCTAA AATAGCTACA 28800 AAAAATTGAG CCATTTAATG CTTATTTCTT ATACCGATGA TATTTCCATA TTCATCAGTT 29400 TGTTTTCCGA GAGCCATCAA AGCTTCGATA AGGTCGATAA TTCCAGGAAT AAAGGTAATA 30000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCTCCA 30000	ATGTCACCTA	TCTTCAATAC	GGTCTGGGAG	CTAAAGCTTA	TCTCAGTGCG	ATTAAAGACC	2340
TTGAGGTAGT CAATATACTT CCAAAGAATA CCGTTATATC ATACAACAAG CTGGTCTGAC 2520 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAATGCA CCAACTGAAA GTTTCTTTGG 2580 GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTGGTCAA 2640 TGATGTGGCA CGTTATATCG AATTCTACAA CACACAACGT TATCAATCAA AATTAAACAA 2700 CCTGACTCCT CTAGAATTCA GGAATCAGGT TGCATAACTT ATCTTTATT ATTTGACTGT 2760 CTACTTGACA GGGAGCCGTT CAGATTGCTT AACCTTTCTA AATTTGCTAA AATAGCTACA 2820 AGAAAACGAG CCATTTAATG CTTATTTCTT ATACCGATGA TATTTCCATA TTCATCAGTT 2940 TGTTTTCCGA GAGCCATCAA AGCTTCGATA AGGTCGATAA TTCCAGGAAT AAAGGTAATA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTCCTG CGTAAAATTT ATGCGCTCCA 3060	TGTATAACGG	TTCTATTATC	GCTTATGAGA	TTAGTCACAA	CAATGAAATC	CACTTGTTAT	2400
CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAATGCA CCAACTGAAA GTTTCTTTGG 2580 GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTGGTCAA 2640 TGATGTGGCA CGTTATATCG AATTCTACAA CACACAACGT TATCAATCAA AATTAAACAA 2700 CCTGACTCCT CTAGAATTCA GGAATCAGGT TGCATAACTT ATCTTTTATT ATTTGACTGT 2760 CTACTTGACA GGGAGCCGTT CAGATTGCTT AACCTTTCTA AATTTGCTAA AATAGCTACA 2880 AAAAAATGAG CCATTTAATG CTTATTTCTT ATACTGTCTT GCCTCACGCT CTCCTCGACC 2880 CTGTTTTCCGA GAGCCATCAA AGCTTCCATA TTCATCAGTT 2940 CTGTTTCCGA GAGCCATCAA AGCTTCGATA AGGTCGATAA TTCCAGGAAT AAAGGTAATA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTCCTG CGTAAAATTT ATGCGCTCCA 3060 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTCCTG CGTAAAAATTT ATGCGCTCCA 3060 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCTCCA 3060 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCTCCA 3060 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCTCCA 3060 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3060 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3060 CTAAAAAATAA TATATAAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCAA 3060 CTAAAAATAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCAA 3060 CTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCAA 3	GAAGACCATT	AAAAAGGGGC	TAGAGCTCAA	TCCAGGAGCC	ACACCTATCA	TCCATAGCGA	2460
GTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTGGTCAA 2640 TGATGTGGCA CGTTATATCG AATTCTACAA CACACAACGT TATCAATCAA AATTAAACAA 2700 CCTGACTCCT CTAGAATTCA GGAATCAGGT TGCATAACTT ATCTTTTATT ATTTGACTGT 2760 CTACTTGACA GGGAGCCGTT CAGATTGCTT AACCTTTCTA AATTTGCTAA AATAGCTACA 2820 AAAAAATGAG CCTTATTATT CTTTTTCTT ATACTGTCTT GCCTCACGCT CTCCTCGACC 2880 CTGTTTCCGA GAGCCATCAA AGCTTCCAT TAAACGATGA TATTTCCATA TTCATCAGTT 2940 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTCCTG CGTAAAAATTT ATGCGCTCCA 3060 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3060 CTAAAAAATAA TATATAAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3060 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3060 CTAAAAAATAA TATATAAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAAATAA TATATAAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAATTA ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA ATGTTTTTTTTTTTTTT	TTGAGGTAGT	CAATATACTT	CCAAAGAATA	CCGTTATATC	ATACAACAAG	CTGGTCTGAC	2520
TGATGTGCA CGTTATATCG AATTCTACAA CACACACGT TATCAATCAA AATTAAACAA 270 CCTGACTCCT CTAGAATTCA GGAATCAGGT TGCATAACTT ATCTTTATT ATTTGACTGT 276 CCTACTTGACA GGGAGCCGTT CAGATTGCTT AACCTTTCTA AATTTGCTAA AATAGCTACA 282 CAAAAAACGAG CCATTTAATG CTTATTTCTT ATACTGTCTT GCCTCACGCT CTCCTCGACC 288 CAAAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 294 CCTAAAAAATAA GAGCCATCAA AGCTTCGATA AGGTCGATAA TTCCAGGAAT AAAGGTAATA 300 CCTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTCCTG CGTAAAAATTT ATGCGCTCCA 306 CCTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 306 CCTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 306 CCTAAAAATAA TATATAAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATTT ATGCGCTCCA 306 CCTAAAAATTT ATGCGCTCCA 306 CCTAAAAATTA 306 CCTAAAAATTT ATGCGCTCCA 306 CCTAAAAATTT ATGCGCTCAAAATTT ATGCGCTCAAAATTT ATGCGCTCAAAATTT ATGCGCTCAAAATTT ATGCGCTCAAAATTT ATGCGCTCAAAATTT ATGCGCTCAAAATTT ATGCGCTCAAAAATTT ATGCGCTCAAAATTT ATGCGCTCAAAATTT ATGCGCTCAAAATTT ATGCGCTCAAAATTT ATGCGCTCAAAATTTAAAA	CTTATCCATG	TCCCGGATTG	GCAAATGTAT	TGATAATGCA	CCAACTGAAA	GTTTCTTTGG	2580
TGATGTGGCA CGTTATATCG AATTCTACAA CACACAACGT TATCAATCAA AATTAAACAA 2700 CCTGACTCCT CTAGAATTCA GGAATCAGGT TGCATAACTT ATCTTTTATT ATTTGACTGT 2760 CTACTTGACA GGGAGCCGTT CAGATTGCTT AACCTTTCTA AATTTGCTAA AATAGCTACA 2820 AGAAAACGAG CCATTTAATG CTTATTTCTT ATACTGTCTT GCCTCACGCT CTCCTCGACC 2880 AAAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 2940 CTAAAAAATAA AACCTGGCCT ATTTTCCTG CGTAAAATTT ATGCGCTCCA 3060 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTCCTG CGTAAAATTT ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCCA 3060 CTAAAATTA ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAATTT ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCCA 3060 CTAAAATTA ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAAATTT ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAATTA ATGCGCTCCA 3060 CTAAAATTT ATGCGCTCCA 3060 CTAAAAATTA ATGCGCTCCA 3060 CTAAAATTA ATGCGCTCCA 3060 CTAAAATTA ATGCGCTCCA 3060 CTAAAATTA ATGCGCTCCA 3	GTTTTTCAAG	ACTGAGTCTT	ACCACCTTAA	GAAATACAAC	TCTTATGATG	AGTTGGTCAA	2640
CTACTTGACA GGGAGCCGTT CAGATTGCTT AACCTTTCTA AATTTGCTAA AATAGCTACA 2820 AGAAAACGAG CCATTTAATG CTTATTTCTT ATACTGTCTT GCCTCACGCT CTCCTCGACC 2880 AAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 2940 TGTTTTCCGA GAGCCATCAA AGCTTCGATA AGGTCGATAA TTCCAGGAAT AAAGGTAATA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTCCTG CGTAAAATTT ATGCGCTCCA 3060	TGATGTGGCA	CGTTATATCG	AATTCTACAA	CACACAACGT	TATCAATCAA	AATTAAACAA	2700
AGAAAACGAG CCATTTAATG CTTATTTCTT ATACTGTCTT GCCTCACGCT CTCCTCGACC 2880 AAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 2940 TGTTTTCCGA GAGCCATCAA AGCTTCGATA AGGTCGATAA TTCCAGGAAT AAAGGTAATA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTCCTG CGTAAAATTT ATGCGCTCCA 3060	CCTGACTCCT	CTAGAATTCA	GGAATCAGGT	TGCATAACTT	ATCTTTTATT	ATTTGACTGT	2760
AAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCCATA TTCATCAGTT 2940 TGTTTTCCGA GAGCCATCAA AGCTTCGATA AGGTCGATAA TTCCAGGAAT AAAGGTAATA 3000 CTAAAAAATAA TATATAAAAA AACCTGGCCT ATTTTCCTG CGTAAAATTT ATGCGCTCCA 3060	CTACTTGACA	GGGAGCCGTT	CAGATTGCTT	AACCTTTCTA	AATTTGCTAA	AATAGCTACA	2820
TGTTTTCCGA GAGCCATCAA AGCTTCGATA AGGTCGATAA TTCCAGGAAT AAAGGTAATA 3000 CTAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCTCCA 3060	AGAAAACGAG	CCATTTAATG	CTTATTTCTT	ATACTGTCTT	GCCTCACGCT	CTCCTCGACC	2880
CTAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAATTT ATGCGCTCCA 3060	AAAAATTGAG	CGTGAGGCTT	TTTGTTTCAT	TAAACGATGA	TATTTCCATA	TTCATCAGTT	2940
	TGTTTTCCGA	GAGCCATCAA	AGCTTCGATA	AGGTCGATAA	TTCCAGGAAT	AAAGGTAATA	3000
ATGCCGCCCA AAAGAACGTT AATAAAACAT AAACTACTAT GTTAGCATAA GACTTTATTT 3120	СТАААААТАА	TATATAAAAA	AACCTGGCCT	ATTTTTCCTG	CGTAAAATTT	ATGCGCTCCA	3060
	ATGCCGCCCA	AAAGAACGTT	AATAAAACAT	AAACTACTAT	GTTAGCATAA	GACTTTATTT	3120

TTACAACTGA	ATTTCATATA	AATGGATTAG	AGTAAGGGA	r aaaagaaati	AGCATAGCTC	3180
TTTTGAAAAT	AAAAAATTA	ATATAATATG	GAAAAAATT	TATTTCATA	ACGTTTCATA	3240
AAAGGTATGT	AATCTAGTAT	TTAGGCAACA	CTATTTTGT	CACTGGTGTCT	AGTAACTTAT	3300
AGATTGATAA	TTTTACTAGT	AAACGTAATT	CTTCGCTTT	A AGAGTTAAAT	GTCTATTTAT	3360
TGTAAGCTAA	ATTGGGAGGT	GAACTTATGT	AAAATTAGAT	AGGTACTGTC	AAGTACGGGA	3420
TGATTATTGA	AACAGCCAGT	ATGCATCATA	AAATCTGTAT	TGCTTAATAA	CTATTTCCTT	3480
AACCAGACAT	CAGTTCATTG	TTTATCATCG	CTACCCTAAG	TCTAGTTTTT	TCAATAGAGC	3540
ATTAGGTAGT	TTTTGATAAT	AAAACTATAT	AAACATGAGA	ATTAGATTTC	GTATTGCATT	3600
CTTCATAATG	AGTTATTTGA	GATTTTCCTT	TGAATAAATA	GATACGAAAT	TCAGTAACTT	3660
CATATATAAA	CGGCTCTATC	ATTGAGATAG	TTTGTCAAAT	GAAGAAATTT	TTAATGGAAA	3720
TAGTTTTAAA	AACATTAGTT	GTAGGCGATG	TAAAAATATT	AATCCAGTGG	ATGCAATAGT	3780
TGCGGAGTAA	AAATAGAGAG	GAGTAATTAG	GAAGTGATAA	AAAATGCTAT	AGCATATATT	3840
ACCAGAAAAA	AAAATAGAAC	ACTTATTATA	TTTGCTATTT	TAACAATTGT	TCTTTCTTGC	3900
TTGTATTCAT	GTTTAACAAT	AATGAAATCA M K S	AGTAATGAAA S N E I		TTTATATGAA L Y E	3960
AGTTCTAATT S S N S	CTTCAATATC S I S	AATTACAAAA I T K	AAAGATGGTA K D G K	AATATTTTAA Y F N	TATTAATCAA I N Q	4020
TTTAAGAATA F K N I	TTGAAAAAAT E K I	AAAAGAGGTT K E V	GAAGAAAAA E E K I	TATTTCAATA F Q Y	TGATGGATTA D G L	4080
GCAAAATTGA A K L K	AAGATCTTAA D L K	AGTAGTTAGT V V S	GGTGAGCAAA G E Q S	GTATAAATAG I N R	AGAAGATTTA E D L	4140
TCTGACGAAT S D E F	TTAAAAATGT K N V	TGTTTCACTA V S L	GAAGCTACAA E A T S	GTAATACTAA N T K	AAGAAATCTT R N L	4200
TTATTTAGTA L F S S	GTGGAGTATT G V F	TAGTTTTAAA S F K	GAAGGAAAAA E G K N	ATATAGAAGA I E E	AAATGATAAG N D K	4260
AATTCAATTC N S I L	TTGTTCATGA V H E	AGAATTTGCT E F A	AAACAAAACA K Q N K	AACTAAAATT L K L	GGGTGATGAA G D E	4320
ATTGATCTTG . I D L E	AATTACTAGA L L D	TACGGAAAAA T E K	AGTGGAAAAA S G K I	TAAAAAGTCA K S H	TAAATTTAAA K F K	4380
ATTATAGGAA '	TCTTTTCTGG F S G	TAAAAAACAG K K Q	GAAACATATA E T Y T	CAGGATTATC G L S	ATCTGATTTT S D F	4440
AGCGAAAATA ' S E N M	rggtttttgt . V F V	AGATTATTCA D Y S	ACTAGCCAAG T S Q E	AAATATTAAA I L N	TAAATCAGAG K S E	4500
AATAATAGAA 1 N N R I	ITGCAAATAA A A N K		TATTCTGGTA Y S G S	GTTTAGAATC L E S	TACAGAGCTT T E L	4560
GCCTTAAACA A	AATTGAAAGA (CTTTAAAATT	GATAAGTCAA	AGTATTCTAT	TAAGAAAGAT	4620

A L N K	L K D	F K I	D K S K Y S I K K D	
AATAAAGCAT N K A F	TCGAAGAGTC E E S	TTTAGAGTCA L E S	GTGAGTGGAA TAAAACATAT AATTAAAATA V S G I K H I I K I	4680
ATGACTTATT M T Y S	CGATTATGTT I M L	AGGTGGAATA G G I	GTTGTTCTTT CATTAATCTT GATTCTATGG V V L S L I L I L W	4740
TTAAGAGAAA L R E R	GAATTTATGA I Y E	AATAGGTATA I G I	TTTTTATCTA TTGGAACAAC TAAGATACAA F L S I G T T K I Q	4800
ATTATAAGGC I I R Q	AATTTATATT F I F	TGAGTTAATA E L I	TTCATATCAA TACCAAGTAT AATATCCTCC F I S I P S I I S S	4860
TTATTTTAG L F L G	GGAATCTACT N L L	ATTAAAAGTA L K V	ATTGTAGAAG GATTTATTAA CTCAGAGAAC I V E G F I N S E N	4920
TCAATGATTT S M I F	TCGGTGGAAG G G S	TTTAATAAAT L I N	AAAAGCAGTT TTATGTTAAA CATAACAACA K S S F M L N I T T	4980
CTTGCAGAAA L A E S	GTTATTTAAT Y L I	ATTAATAAGT L I S	ATTATTGTTT TATCAGTTGT AATGGCCTCT I I V L S V V M A S	5040
TCATTAATAT S L I L	F K K	PQE	ATATTATCAA AAATAAGTTA GGAGCAAATA I L S K I S .	5100
ATGGATATAT M D I L	TAGAAATAAA E I K	GAATGTAAAT N V N	TACAGTTACG CAAATTCTAA AGAAAAAGTT Y S Y A N S K E K V	5160
L S G V	N Q K	F E L	GGAAAGTTTT ATGCGATAGT AGGGAAGTCA G K F Y A I V G K S	5220
GGAACAGGAA G T G K	S T L	L S L	CTTGCAGGAC TTGATAAAGT TCAAACAGGA L A G L D K V Q T G	5280
AAAATCTTGT K I L F	K N E	DIE	K K G Y S N H R K N	5340
AATATATCTT N I S L	V F Q	N Y N	TTAATAGATT ATTTATCGCC GATTGAAAAT L I D Y L S P I E N	5400
ATTAGACTAG I R L V	N K S	V D E	SILFELGLDK	5460
K Q I K	R N V	M K L	S G G Q Q R V A I	5520
A R A L	V S D	API	ATACTAGCTG ATGAGCCTAC CGGTAACCTA I L A D E P T G N L	5580
GACAGTGTTA D S V T	CTGCTGGAGA A G E	AATAATT ((SEQ ID NO:27)	5607

FIG. 5a

TÖSTIEGÖTK	SUIPERRAGU	TVRVHAKVVE	GTRERIQIFE	GVVISRKGQG	50
ISEMYTVRKI	SGGIGVERTF	PIHTPRVDKI	EVVRYGKVRR	AKLYYLRALQ	100
GKAARIKEIR	R (SEQ ID	NO:28)			111
		FIG. 5	b		
				KSSIRNIPID	50
NDTLFFLHEF	TKNKNDRLFD	KLSNNAVNKT	IRKITGREVR	VHSLRHTFAS	100
YLISISQVLD	HENLNITLEV	YAHQLQEQKD	RNDKLNQRNL	GQNSSKPLFT	150
CNEYVPCRNR	TSNYSLGGSC	YIH (SEQ	ID NO:29)		173
		FIG. 5	C		
VVICANIE TENIN					
				KEVEEKIFQY	50
			VSLEATSNTK		100
			GDEIDLELLD		150
KFKIIGIFSG	KKQETYTGLS	SDFSENMVFV	DYSTSQEILN	KSENNRIANK	200
ILMYSGSLES	TELALNKLKD	FKIDKSKYSI	KKDNKAFEES	LESVSGIKHI	250
IKIMTYSIML	GGIVVLSLIL	ILWLRERIYE	IGIFLSIGTT	KIQIIRQFIF	300
ELIFISIPSI	ISSLFLGNLL	LKVIVEGFIN	SENSMIFGGS	LINKSSFMLN	350
ITTLAESYLI	LISIIVLSVV	MASSLILFKK	PQEILSKIS		389
(SEQ ID NO:	30)				
		FIG. 5	d		
			GKFYAIVGKS		50
			NISLVFQNYN		100
IRLVNKSVDE	SILFELGLDK	KQIKRNVMKL	SGGQQQRVAI	ARALVSDAPI	150
ILADEPTGNL	DSVTAGEII	(SEQ ID NO:	31)		169

FIG. 5e

CATATGACAA	TATTTTTCAA	AGTCTACATC	ACTTACTCGC	CTGTCGTGGA	AAATCTGGCA	60
ATACATTAAT	CGACCAATTA	GTTGCTGATG	GTTTACTTCA	TGCAGATAAT	CACTACCATT	120
TTTTCAATGG	GAAGTCTCTG	GCCACTTTCA	ATACTAACCA	ATTGATTCGC	GAAGTTGTCT	180
ATGTTGAAAT	ATCCTTAGAT	ACTATGTCTA	GTGGTGAACA	TGATTTAGTA	AAAGTTAACA	240
TTATCAGACC	CACTACCGAG	CATACTATCC	CCACGATGAT	GACAGCTAGC	CCCTATCATC	300
AAGGTATCAA	TGATCCTGCC	GCAGACCAAA	AAACATACCA	AATGGAGGGT	GCGCTAGCAG	360
TTAAACAGCC	TAAACACATA	CAAGTTGACA	CAAAACCATT	TAAAGAAGAA	GTAAAACATC	420
CTTCAAAATT	ACCCATCAGC	CCTGCAACTG	AAAGCTTCAC	ACACATTGAC	AGTTATAGTC	480
TCAATGACTA	TTTTCTTTCT	CGTGGTTTTG	CTAATATATA	CGTTTCAGGT	GTGGGTACTG	540
CTGGCTCTAC	GGGTTTCATG	ACCAGTGGGG	ATTACCAACA	AATACAAAGC	TTTAAAGCAG	600
TCATTGATTG	GTTAAATGGT	AAGGTTACTG	CATTCACAAG	TCATAAACGA	GATAAACAAG	660
TCAAGGCTGA	TTGGTCAAAC	GGCCTTGTAG	CAACCACAGG	TAAATCTTAT	CTCGGTACCA	720
TGTCAACTGG	TTTAGCAACA	ACTGGCGTTG	AGGGGCTGAA	AGTCATTATC	GCTGAAGCCG	780
CAATCTCCAC	ATGGTATGAT	TATTATCGAG	AAAATGGGCT	TGTGTGTAGT	CCAGGCGGCT	840
ACCCCGGTGA	AGATTTAGAC	GTTTTAACAG	AATTAACATA	CTCACGAAAC	CTCTTAGCTG	900
GTGATTACAT	CAAAAACAAC	GATTGCTATC	AAGCATTGTT	AAATGAACAA	TCAAAAGCAA	960
TTGACCGTCA	AAGTGGGGAT	TACAACCAAT	ACTGGCATGA	CCGTAATTAC	CTAACTCACG	1020
TCAATAATGT	CAAAAGTCGA	GTAGTTTACA	CTCATGGACT	ACAGGATTGG	AATGTTAAGC	1080
CAAGACATGT	CTACAAAGTT	TTCAATGCAT	TGCCTCAAAC	CATCAAAAAA	CACCTTTTTT	1140
TACATCAAGG	TCAACATGTG	TATATGCATA	ATTGGCAGTC	GATTGATTTT	CGTGAAAGCA	1200
TGAATGCCTT	ACTAAGCCAA	GAACTACTTG	GCATTGACAA	TCATTTCCAA	TTAGAAGAGG	1260
TCATTTGGCA	AGATAATACT	ACTGAGCAAA	CTTGGCAAGT	TTTAGATGCT	TTCGGAGGAA	1320
ACCATCAAGA	GCAAATTGGT	TTAGGTGATA	GTAAAAAACT	TATTGATAAC	CATTATGACA	1380
AAGAAGCCTT	TGATACTTAT	TGTAAAGACT	TCAATGTGTT	CAAAAATGAT	CTTTTCAAGG	1440
GAAATAATAA	AACCAATCAA	ATCACTATTA	ATCTTCCTCT	AAAGAAAAAT	TATCTCCTGA	1500
ATGGACAGTG	CAAACTCCAT	CTACGTGTTA	AAACTAGTGA	CAAAAAGGCC	ATTTTATCAG	1560
CCCAAATCTT	AGACTATGGT	CCTAAAAAAC	GATTCAAAGA	TACACCAACC	ATCAAATTCT	1620
TAAACAGCCT	TGATAATGGT	AAAAATTTTG	CCAGAGAAGC	TTTACGTGAA	CTCCCGTTTA	1680
CTAAAGATCA	TTATCGTGTC	ATCAGTAAAG	GTGTCTTGAA	CCTTCAAAAT	CGTACAGACT	1740
TACTTACAAT	TGAGGCTATC	GAGCCAGAAC	AATGGTTTGA	TATCGAGTTT	AGCCTCCAAC	1800
	TCAATTGAGT					1860
TTGAACATAC	CATTCGAGAT	AATGCTAGTT	ACTCTATAAC	AGTAGATTTG	AGTCAATCTT	1920
ATTTAACTAT	CCCAACTAAT	CAAGGAAATT	AACTTATGAA	ACTTCTTACT	AAAGAACGGT	1980
TTGATGATTC	TCAACACTTT	TGGTACCAGA	TCAATTTATT	ACAAGAGAGT	AACTTCGGAG	2040
	CCATGATAAT					2100
TACAAGGTTC	CGGAAGTTCG	AATCATTTCT	GGTATTTTGG	CAATACTACT	GATACTTCCA	2160
TCCTTATGAT	TGCTCATTTA	AATCGAAAAT	TCTATATTCA	GGTTAATTTA	AAGGACTTTG	2220
	CAATTTAATA					2280
	CGATACCCTA					2340
AACGCGAGGG	AGACTGATTA	ATGTCATCTT	ATTGGAATAA	CTATCCTGAA	СТТАААААА	2400

	ATATTGATGA	AACCAATCAA	CTAATTCAAG	AAAGAATACA	GGTCAGAAAT	AAAGATATTG	2460
	AAGCGGCGCT	AAGCCAACTC	ACAGCTGCGG	GAGGAAAACA	GCTCAGACCA	GCATTCTTTT	2520
	ACCTTTTTTC	TCAACTTGGT	AATAAGGAGA	ATCAAGATAC	TCAGCAACTA	AAGAAAATCG	2580
	CTGCTTCTTT	AGAAATCCTT	CACGTTGCTA	CATTAATCCA	TGATGATGTC	ATTGATGACT	2640
	CACCACTAAG	ACGTGGAAAT	ATGACCATTC	AAAGCAAGTT	TGGCAAAGAC	ATCGCAGTTT	2700
	ATACTGGGGA	TTTACTTTTC	ACAGTCTTTT	TCGATCTTAT	TTTAGAATCT	ATGACTGATA	2760
	CACCATTTAT	GAGGATTAAT	GCAAAATCTA	TGCGTAAAAT	TCTCATGGGA	GAATTGGACC	2820
	AGATGCACCT	TCGTTACAAT	CAACAACAAG	GTATCCATCA	CTATTTACGT	GCGATTTCAG	2880
	GTAAGACAGC	CGAACTCTTT	AAATTAGCTA	GCAAAGAAGG	AGCTTACTTT	GGTGGTGCAG	2940
	AGAAGGAGGT	TGTTCGTCTA	GCAGGCCATA	TCGGCTTTAA	CATTGGTATG	ACATTCCAAA	3000
	TTTTGGATGA	TATCCTGGAT	TATACTGCAG	ATAAAAAAAC	ATTTAATAAG	CCTGTCTTAG	3060
	AGGATTTAAC	ACAAGGCGTT	TACAGCCTTC	CTCTACTTCT	TGCCATTGAA	GAAAATCCTG	3120
	ATATTTTCAA	ACCTATTTTA	GATAAAAAA	CAGATATGGC	TACTGAAGAC	ATGGAAAAA	3180
	TTGCTTATCT	CGTCGTTTCC	CATAGAGGTG	TTGACAAAGC	TCGCCATCTA	GCTCGTAAAT	3240
		AGCTATTAGT					3300
	TGCTACAATT	AACTAATTAC	CTTTTAAAAC	GCAAAATTTA	AATAATAAA	AAACATTCCA	3360
	CAATGCTAGA	AAAGCAGTTA	GGGAATGTTT	TTTTATTATC	ATTTATTTAT	CGCACCTATC	3420
	AATCATCATA	GATCACCATC	ATCAGCGGCT	TTCAGCTGAC	GGTAACGTTG	ACTACTTTGA	3480
	GACAATTCTT	GAGGAGAACC	TTCCAACTCT	AATTGCCCAT	TTTCTATAAA	TAAGATACGA	3540
	TCAGCATGTT	CAATACCTTT	TAAGTGATGT	GTAATCCAAA	CTAAGGTCTT	ACCTTCCAAT	3600
	TCTTTCATAA	ATACCCTTAG	TAAGGCTTGT	TCAGTAATAG	GATCAAGTCC	AACAGTTGGC	3660
	TCATCTAAGA	TAACAATTGG	GACATCTTTT	AGTAAGATTC	TAGCCAAAGC	AATTCTATGC	3720
	CTTTCGCCAC	CTGAAAACCT	AAGTCCAGCT	TCATCAACCA	TTGTATAGAG	ACCATCTGAT	3780
	AAATCAGTGA	CCATCTCTTT	CAATCCAACT	CGTTCAAGAA	CTTTCCATAC	ATCTTCTTCA	3840
	CTAGCATCTT	GGTTTCCAAT	GCGAATGTTA	TTTAGCAGGG	TTGTATTAAA	AAGGTAGGGC	3900
	GCTTGTTGTA	TCACTCCAAT	ATAGTTAGAA	ATGCAATCAC	CAACTATTGA	AACATCAGCA	3960
	CCGCCTAGGG	TAATCTTCCC	TTGACTTGCT	TTCAAGTCGC	CACGAAGTAG	ACTAGCTAAG	4020
	GTACTCTTGC	CAGAACCACT	CCGCCCTAAA	ATAGCAATTT	TTTCTCCTTC	TTTAATATCC	4080
	AAATCTAAAT	GATGCAAAAC	CCATTTCTCT	TGTGGCTTAT	ACTGGAAACT	TAAATTCTTG	4140
,	ACGGAAAAAT	CATATGGCTT	ATTAGGCAAT	T (SEQ ID	NO:32)		4171

FIG. 6a

YUI	NIFQSLHH	LLACRGKSGN	TLIDQLVADG	LLHADNHYHF	FNGKSLATFN	50
TNO	QLIREVVY	VEISLDTMSS	GEHDLVKVNI	IRPTTEHTIP	TMMTASPYHQ	100
GIN	NDPAADQK	TYQMEGALAV	KQPKHIQVDT	KPFKEEVKHP	SKLPISPATE	150
SFT	THIDSYSL	NDYFLSRGFA	NIYVSGVGTA	GSTGFMTSGD	YQQIQSFKAV	200
IDV	VLNGKVTA	FTSHKRDKQV	KADWSNGLVA	TTGKSYLGTM	STGLATTGVE	250
GLF	KVIIAEAA	ISTWYDYYRE	NGLVCSPGGY	PGEDLDVLTE	LTYSRNLLAG	300
DYI	KNNDCYQ	ALLNEQSKAI	DRQSGDYNQY	WHDRNYLTHV	NNVKSRVVYT	350
HGI	LQDWNVKP	RHVYKVFNAL	PQTIKKHLFL	HQGQHVYMHN	WQSIDFRESM	400
NAI	LLSQELLG	IDNHFQLEEV	IWQDNTTEQT	WQVLDAFGGN	HQEQIGLGDS	450
KKI	TIDNHYDK	EAFDTYCKDF	NVFKNDLFKG	NNKTNQITIN	LPLKKNYLLN	500
GQC	KLHLRVK	TSDKKAILSA	QILDYGPKKR	FKDTPTIKFL	NSLDNGKNFA	550
REF	ALRELPFT	KDHYRVISKG	VLNLQNRTDL	LTIEAIEPEQ	WFDIEFSLQP	600
SIY	QLSKGDN	LRIILYTTDF	EHTIRDNASY	SITVDLSQSY	LTIPTNQGN	649
(SE	Q ID NO:	:33)				
			TOTAL C	L.		

FIG. 6b

MKLLTKERFD	DSQHFWYQIN	LLQESNFGAV	FDHDNKNIPQ	VVATIVDDLQ	50
GSGSSNHFWY	FGNTTDTSIL	MIAHLNRKFY	IQVNLKDFDF	ALNLIAINNW	100
KSLLQTQLEA	LNDTLAIFQ	(SEQ ID NO):34)		119

FIG. 6c

MSSYWNNYPE	LKKNIDETNQ	LIQERIQVRN	KDIEAALSQL	TAAGGKQLRP	50
AFFYLFSQLG	NKENQDTQQL	KKIAASLEIL	HVATLIHDDV	IDDSPLRRGN	100
MTIQSKFGKD	IAVYTGDLLF	TVFFDLILES	MTDTPFMRIN	AKSMRKILMG	150
ELDQMHLRYN	QQQGIHHYLR	AISGKTAELF	KLASKEGAYF	GGAEKEVVRL	200
AGHIGFNIGM	TFQILDDILD	YTADKKTFNK	PVLEDLTQGV	YSLPLLLAIE	250
ENPDIFKPIL	DKKTDMATED	MEKIAYLVVS	HRGVDKARHL	ARKFTEKAIS	300
DINKLPQNSA	KKQLLQLTNY	LLKRKI (SE	EQ ID NO:35)		326

FIG. 6d

LPNKPYDFSV	KNLSFQYKPQ	EKWVLHHLDL	DIKEGEKIAI	LGRSGSGKST	50
LASLLRGDLK	ASQGKITLGG	ADVSIVGDCI	SNYIGVIQQA	PYLFNTTLLN	100
NIRIGNQDAS	EEDVWKVLER	VGLKEMVTDL	SDGLYTMVDE	AGLRFSGGER	150
HRIALARILL	KDVPIVILDE	PTVGLDPITE	QALLRVFMKE	LEGKTLVWIT	200
HHLKGIEHAD	RILFIENGQL	ELEGSPQELS	QSSQRYRQLK	AADDGDL	247
(SEQ ID NO:	:36)				

FIG. 6e

AATTCTATTT	GGAGGTTTTT	CTTGAATAAA	TGGTTAGTTA	AGGCAAGTTC	CTTAGTTGTT	60
TTAGGTGGTA	TGGTTTTATC	TGCGGGTTCC	CGAGTTTTAG	CGGATACTTA	TGTCCGTCCA	120
ATTGATAATG	GTAGAATTAC	AACAGGTTTC	AATGGTTATC	CTGGACATTG	TGGGGTGGAT	180
TATGCTGTTC	CGACTGGAAC	GATTATTAGG	GCAGTGGCAG	ATGGTACTGT	GAAATTTGCA	240
GGAGCTGGAG	CCAACTTTTC	TTGGATGACA	GACTTAGCAG	GAAATTGTGT	CATGATTCAA	300
CATGCGGATG	GAATGCATAG	TGGTTACGCT	CATATGTCAC	GTGTGGTGGC	TAGGACTGGG	360
GAAAAAGTCA	AACAAGGAGA	TATCATCGGT	TACGTAGGAG	CAACTGGTAT	GGCGACGGGA	420
CCTCACCTTC	ATTTTGAATT	TTTACCAGCT	AACCCTAATT	TTCAAAATGG	TTTCCATGGA	480
CGTATCAATC	CAACGTCACT	AATTGCTAAC	GTTGCGACCT	TTAGTGGAAA	AACGCAAGCA	540
TCAGCTCCAA	GCATTAAGCC	ATTACAATCA	GCTCCTGTAC	AGAATCAATC	TAGTAAATTA	600
AAAGTGTATC	GAGTAGATGA	ATTACAAAAG	GTTAATGGTG	TTTGGTTAGT	CAAAAATAAC	660
ACCCTAACGC	CGACTGGGTT	TGATTGGAAC	GATAATGGTA	TACCAGCATC	AGAAATTGAT	720
GAGGTTGATG	CTAATGGTAA	TTTGACAGCT	GACCAGGTTC	TTCAAAAAGG	TGGTTACTTT	780
ATCTTTAATC	CTAAAACTCT	TAAGACTGTA	GAAAAACCCA	TCCAAGGAAC	AGCTGGTTTA	840
ACTTGGGCTA	AGACACGCTT	TGCTAATGGT	AGTTCAGTTT	GGCTTCGCGT	TGACAACAGT	900
CAAGAACTGC				GTTTTAAATG		960
TACTAACTAA	GTACAATTTC	TTTAAACCGT	CTGAAAATAA	TTTTATAGTC	CAGTAAAGTG	1020
				TGAAGCAATG		1080
AAAAGGTACT	ATTGACATCG	ACAATGGCAG	CTTCGCTATT	ATCAGTCGCA	AGTGTTCAAG	1140
CACAAGAAAC	AGATACGACG	TGGACAGCAC	GTACTGTTTC	AGAGGTAAAG	GCTGATTTGG	1200
TAAAGCAAGA	CAATAAATCA	TCATATACTG	TGAAATATGG	TGATACACTA	AGCGTTATTT	1260
CAGAAGCAAT	GTCAATTGAT	ATGAATGTCT	TAGCAAAAAT	TAATAACATT	GCAGATATCA	1320
				TCAGAAGAGT	CATACTGCCA	1380
	AATAGAAACA					1440
				TTCTCTCAAT		1500
				AATGAAGACA		1560
			-	AGCTGTTAGT		1620
				TTCAGAAGTT		1680
				AACAACAGTA		1740
				ACCGGTAAGA		1800
				AGAAACTGGT		1860
				AACAGCTACA		1920
				AGCTCCAACA		1980
				TCCTGAAAAT		2040
				TTATGGAGTT		2100
				TTTAGCAGTC		2160
				CTCTACACAA		2220
				CTCAAATACA		2280
				TGGCGTTACT		2340
ATGACCATGT	TCACGTATCA	TTTAACAAAT	AATATAAAAA	AGGAAGCTAT	TTGGCTTCTT	2400

TTTTATATGC CTTGAATAGA	CTTTCAAGGT	TCTTATCTAA	TTTTTATTAA	ATTGAGGAGA	2460
TTAAGCTATA AGTCTGAAAC	TACTTTCACG	TTAACCGTGA	CTAAATCAAA	ACGTTAAAAC	2520
TAAAATCTAA GTCTGTAAAG	ATTATTGAAA	ACGCTTTAAA	AACAGATATA	ATAAGGTTTG	2580
TAGATATCTA AAATTAAAAA	AGATAAGGAA	GTGAGAATAT	GCCACATCTA	AGTAAAGAAG	2640
CTTTTAAAAA GCAAATAAAA	AATGGCATTA	TTGTGTCATG	TCAAGCTTTG	CCTGGGGAGC	2700
CTCTTTATAC TGAAAGTGGA	GGTGTTATGC	CTCTTTTAGC	TTTGGCAGCT	CAAGAAGCAG	2760
GAGCGGTTGG TATAAGAGCC	AATAGTGTCC	GCGACATTAA	GGAAATTCAA	GAAGTTACTA	2820
ATTTACCTAT CATCGGCATT	ATTAAACGTG	AATATCCTCC	ACAAGAACCA	TTTATCACTG	2880
CTACGATGAC AGAGGTGGAT	CAATTAGCTA	GTTTAGATAT	TGCAGTAATA	GCCTTAGATT	2940
GTACACTTAG AGAGCGTCAT	GATGGTTTGA	GTGTAGCTGA	GTTTATTCAA	AAGATAAAAG	3000
GGAAATATCC TGAACAGTTG	CTAATGGCTG	ATATAAGTAC	TTTTGAAGAA	GGTAAAAATG	3060
CTTTTGAAGC AGGAGTTGAT	TTTGTGGGTA	CAACTCTATC	TGGATACACA	GATTACAGCC	3120
GCCAAGAAGA AGGACCGGAT	ATAGAACTCC	TTAATAAGCT	TTGTCAAGCC	GGTATAGATG	3180
TGATTGCGGA AGGTAAAATT	CATACTCCTA	AGCAAGCTAA	TGAAATTAAT	CATATAGGTG	3240
TTGCAGGAAT TGTAGTTGGT	GGTGCTATCA	CTAGACCAAA	AGAAATAGCG	GAGCGTTTCA	3300
TCTCAGGACT TAGTTAAAAG	TGTTACTCAA	AAATCAAAAT	CAAAATAAAA	AAGGGGAATA	3360
GTTATGAGTA TCAAAAAAG	TGTGATTGGT	TTTTGCCTCG	GAGCTGCAGC	ATTATCAATG	3420
TTTGCTTGTG TAGACAGTAG	TCAATCTGTT	ATGGCTGCCG	AGAAGGATAA	AGTCGAAATT	3480
(SEQ ID NO:37)					

FIG. 7a

NSIWRFFLNK	WLVKASSLVV	LGGMVLSAGS	RVLADTYVRP	IDNGRITTGF	50
NGYPGHCGVD	YAVPTGTIIR	AVADGTVKFA	GAGANFSWMT	DLAGNCVMIQ	100
HADGMHSGYA	HMSRVVARTG	EKVKQGDIIG	YVGATGMATG	PHLHFEFLPA	150
NPNFQNGFHG	RINPTSLIAN	VATFSGKTQA	SAPSIKPLQS	APVQNQSSKL	200
KVYRVDELQK	VNGVWLVKNN	TLTPTGFDWN	DNGIPASEID	EVDANGNLTA	250
DQVLQKGGYF	IFNPKTLKTV	EKPIQGTAGL	TWAKTRFANG	SSVWLRVDNS	300
QELLYK (S	SEQ ID NO:38	3)			306

FIG. 7b

MKMNKKVLLT	STMAASLLSV	ASVQAQETDT	TWTARTVSEV	KADLVKQDNK	50
SSYTVKYGDT	LSVISEAMSI	DMNVLAKINN	IADINLIYPE	TTLTVTYDQK	100
SHTATSMKIE	TPATNAAGQT	TATVDLKTNQ	VSVADQKVSL	NTISEGMTPE	150
		KEVLAQEQAV			200
		VSPASVAAET			250
VKVVTPKVET	GASPEHVSAP	AVPVTTTSTA	TDSKLQATEV	KSVPVAQKAP	300
TATPVAQPAS	TTNAVAAHPE	NAGLQPHVAA	YKEKVASTYG	VNEFSTYRAG	350
DPGDHGKGLA	VDFIVGKNQA	LGNEVAQYST	QNMAANNISY	VIWQQKFYSN	400
TNSIYGPANT	WNAMPDRGGV	TANHYDHVHV	SFNK (SEQ	ID NO:39)	434

FIG. 7c

MPHLSKEAFK	KQIKNGIIVS	CQALPGEPLY	TESGGVMPLL	ALAAQEAGAV	50
GIRANSVRDI	KEIQEVTNLP	IIGIIKREYP	PQEPFITATM	TEVDQLASLD	100
IAVIALDCTL	RERHDGLSVA	EFIQKIKGKY	PEQLLMADIS	TFEEGKNAFE	150
AGVDFVGTTL	SGYTDYXRQE	EGPDIELLNK	LCQAGIDVIA	EGKIHTPKQA	200
NEINHIGVAG	IVVGGAITRP	KEIAERFISG	LS (SEQ II	NO:40)	232

FIG. 7d

MSIKKSVIGF CLGAAALSMF ACVDSSQSVM AAEKDKVEI (SEQ ID NO:41)

39

FIG. 7e

3 max 2 3 max	202222222000	3 000 mm 0 3 0 3			
ATGAAAATGA				CAGCTTCGCT	50
ATTATCAGTC	GCAAGTGTTC	AAGCACAAGA	AACAGATACG	ACGTGGACAG	100
CACGTACTGT	TTCAGAGGTA		TGGTAAAGCA	AGACAATAAA	150
TCATCATATA	CTGTGAAATA	TGGTGATACA	CTAAGCGTTA	TTTCAGAAGC	200
AATGTCAATT	GATATGAATG	TCTTAGCAAA	AATTAATAAC	ATTGCAGATA	250
TCAATCTTAT	TTATCCTGAG	ACAACACTGA	CAGTAACTTA		300
AGTCATACTG	CCACTTCAAT	GAAAATAGAA	ACACCAGCAA	CAAATGCTGC	350
TGGTCAAACA	ACAGCTACTG	TGGATTTGAA	AACCAATCAA	GTTTCTGTTG	400
CAGACCAAAA	AGTTTCTCTC	AATACAATTT	CGGAAGGTAT	GACACCAGAA	450
GCAGCAACAA	CGATTGTTTC	GCCAATGAAG	ACATATTCTT	CTGCGCCAGC	500
TTTGAAATCA	AAAGAAGTAT	TAGCACAAGA	GCAAGCTGTT	AGTCAAGCAG	550
CAGCTAATGA	ACAGGTATCA	ACAGCTCCTG	TGAAGTCGAT	TACTTCAGAA	600
GTTCCAGCAG	CTAAAGAGGA	AGTTAAACCA	ACTCAGACGT	CAGTCAGTCA	650
GTCAACAACA	GTATCACCAG	CTTCTGTTGC	CGCTGAAACA	CCAGCTCCAG	700
TAGCTAAAGT	AGCACCGGTA	AGAACTGTAG	CAGCCCCTAG	AGTGGCAAGT	750
GTTAAAGTAG	TCACTCCTAA	AGTAGAAACT	GGTGCATCAC	CAGAGCATGT	800
ATCAGCTCCA	GCAGTTCCTG	TGACTACGAC	TTCAACAGCT	ACAGACAGTA	850
AGTTACAAGC	GACTGAAGTT	AAGAGCGTTC	CGGTAGCACA	AAAAGCTCCA	900
ACAGCAACAC	CGGTAGCACA	ACCAGCTTCA	ACAACAAATG	CAGTAGCTGC	950
ACATCCTGAA	AATGCAGGGC	TCCAACCTCA	TGTTGCAGCT	TATAAAGAAA	1000
AAGTAGCGTC	AACTTATGGA	GTTAATGAAT	TCAGTACATA	CCGTGCAGGT	1050
GATCCAGGTG	ATCATGGTAA	AGGTTTAGCA	GTCGACTTTA	TTGTAGGTAA	1100
AAACCAAGCA	CTTGGTAATG	AAGTTGCACA	GTACTCTACA	CAAAATATGG	1150
CAGCAAATAA	CATTTCATAT	GTTATCTGGC		TTACTCAAAT	1200
ACAAATAGTA	TTTATGGACC	TGCTAATACT	TGGAATGCAA	TGCCAGATCG	1250
TGGTGGCGTT	ACTGCCAACC	ATTATGACCA	TGTTCACGTA	TCATTTAACA	
AATAA			TOTTORCGIA	TCATITAACA	1300
					1305

(SEQ ID NO:42)

FIG. 8

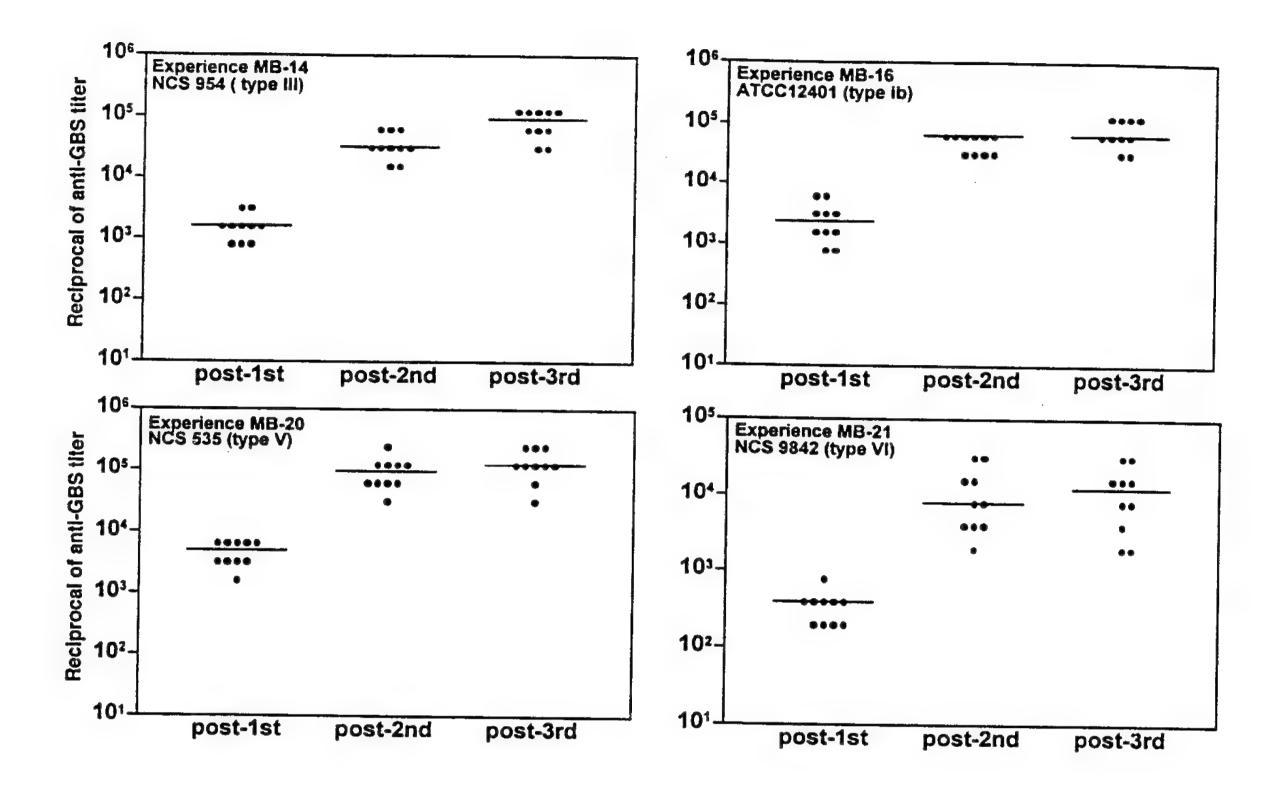
CAAGAAACAG	ATACGACGTG	GACAGCACGT	ACTGTTTCAG	AGGTAAAGGC	50
TGATTTGGTA	AAGCAAGACA	ATAAATCATC	ATATACTGTG		100
ATACACTAAG	CGTTATTTCA	GAAGCAATGT	CAATTGATAT		150
GCAAAAATTA	ATAACATTGC	AGATATCAAT	CTTATTTATC	CTGAGACAAC	200
ACTGACAGTA	ACTTACGATC	AGAAGAGTCA	TACTGCCACT	TCAATGAAAA	
TAGAAACACC	AGCAACAAAT	GCTGCTGGTC	AAACAACAGC	TACTGTGGAT	250
TTGAAAACCA	ATCAAGTŢTC	TGTTGCAGAC	CAAAAAGTTT	CTCTCAATAC	300
AATTTCGGAA	GGTATGACAC	CAGAAGCAGC	AACAACGATT	GTTTCGCCAA	350
TGAAGACATA	TTCTTCTGCG	CCAGCTTTGA	AATCAAAAGA	AGTATTAGCA	400
CAAGAGCAAG	CTGTTAGTCA	AGCAGCAGCT	AATGAACAGG	TATCAACAGC	450 500
TCCTGTGAAG	TCGATTACTT	CAGAAGTTCC	AGCAGCTAAA	GAGGAAGTTA	550
AACCAACTCA	GACGTCAGTC	AGTCAGTCAA	CAACAGTATC	ACCAGCTTCT	600
GTTGCCGCTG	AAACACCAGC	TCCAGTAGCT	AAAGTAGCAC	CGGTAAGAAC	650
TGTAGCAGCC	CCTAGAGTGG	CAAGTGTTAA	AGTAGTCACT	CCTAAAGTAG	700
AAACTGGTGC	ATCACCAGAG	CATGTATCAG	CTCCAGCAGT	TCCTGTGACT	750
ACGACTTCAA	CAGCTACAGA	CAGTAAGTTA	CAAGCGACTG	AAGTTAAGAG	800
CGTTCCGGTA	GCACAAAAAG	CTCCAACAGC	AACACCGGTA	GCACAACCAG	850
CTTCAACAAC	AAATGCAGTA	GCTGCACATC	CTGAAAATGC	AGGGCTCCAA	900
CCTCATGTTG	CAGCTTATAA	AGAAAAAGTA	GCGTCAACTT	ATGGAGTTAA	950
TGAATTCAGT	ACATACCGTG	CAGGTGATCC	AGGTGATCAT	GGTAAAGGTT	1000
TAGCAGTCGA	_	GGTAAAAACC		TAATGAAGTT	1050
GCACAGTACT		TATGGCAGCA	AATAACATTT	CATATGTTAT	1100
CTGGCAACAA		CAAATACAAA	TACTATTAT	GGACCTGCTA	1150
ATACTTGGAA	TGCAATGCCA	GATCGTGGTG	GCGTTACTGC	CAACCATTAT	1200
GACCATGTTC		TAACAAATAA	(SEQ ID		1230
			, 2 5		1230

FIG. 9

QETDTTWTAR	TVSEVKADLV	KQDNKSSYTV	KYGDTLSVIS	EAMSIDMNVL	50
AKINNIADIN	LIYPETTLTV	TYDQKSHTAT	SMKIETPATN	AAGQTTATVD	100
LKTNQVSVAD	QKVSLNTISE	GMTPEAATTI	VSPMKTYSSA	PALKSKEVLA	150
QEQAVSQAAA	NEQVSTAPVK	SITSEVPAAK	EEVKPTQTSV	SQSTTVSPAS	200
VAAETPAPVA	KVAPVRTVAA	PRVASVKVVT	PKVETGASPE	HVSAPAVPVT	250
TTSTATDSKL	QATEVKSVPV	AQKAPTATPV	AQPASTTNAV	AAHPENAGLQ	300
	ASTYGVNEFS				350
AQYSTQNMAA	NNISYVIWQQ	KFYSNTNSIY	GPANTWNAMP	DRGGVTANHY	400
DHVHVSFNK	(SEQ ID NO:				409

FIG. 9a

Fig. 10



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SEQUENCE LISTING

<110> BioChem Vaccins RIOUX, Clément DENIS, Martin BRODEUR, Bernard R. HAMEL, Josée CHARLEBOIS, Isabelle BOYER, Martine <120> NOVEL GROUP B STREPTOCOCCUS ANTIGENS <130> 12806-9PCT <150> 60/075,425 <151> 1998-02-20 <160> 44 <170> FastSEQ for Windows Version 3.0 <210> 1 <211> 4514 <212> DNA <213> Streptococcus <220> <221> CDS <222> (3)...(464) <221> CDS <222> (534)...(887) <223> <221> CDS <222> (1024)...(1767) <221> CDS <222> (1841)...(4288) <221> CDS <222> (2735)...(4288) <400> 1

ta tct ggc aaa gag cca gct aat cgt ttt agt tgg gct aaa aat aaa

1

Ser Gly Lys Glu Pro Ala Asn Arg Phe Ser Trp Ala Lys Asn Lys

10

47

					ttc Phe		_			_	_					95
	_	_			ata Ile	_				_			_			143
					ttg Leu							_				191
_		_			gat Asp		-					_	_			239
					ctt Leu 85				_							287
					ttt Phe								_			335
		_		_	tta Leu	_				_	_	_		-	_	383
				-	atg Met		-				_		_	_		431
					tat Tyr						taat	agaa	aag t	catct	agtga	484
taga	actaa	aca g	gtate	gatat	ig gt	tatgt	caaa	a gta	attta	agga	ggag	gaaga	Me	_	et act er Thr	542
					gca Ala					_	_					590
					tta Leu									_		638
					gaa Glu 195											686
					gta Val											734

				-					-	-	gtt Val 235		tta Leu	782
											gtt Val			830
											ctt Leu			878
tta Leu	taa	tacta	act t	cagco	egtte	cg at	ittag	gttga	a acq	ggcti	tta			927
								atg	act	gag	aac aac Asn	tgg		987 1041
								_	_	_	ggt Gly			1089
											cgc Arg			1137
											att Ile			1185
											acc Thr 340			1233
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	at cac is His														1713
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aat to Asn *	ga aaaa *	agtca	aaa t	cact	gact	et ct	gtga	attaa	a aat	itgta	attt	ttta	atato	ctg	1817
ttttag	gtgct (tatta	attgt	t ga	Me						g Th			et gtt er Val	1870
gag ca	gtgct (aa cta ln Leu	aag	agt	gtt	Me 52 ttt	et I] 20 ggg	le Hi	ls Le	eu Ly tot	/s Ai 52 cca	rg Th	nr Il	le Se	er Val	1870 1918
gag ca Glu Gl 530 tta at	aa cta	aag Lys ctt	agt Ser gtg	gtt Val 535	Me 52 ttt Phe gtt	et I] 20 999 Gly atc	caa Gln gct	tta Leu gtc	tct Ser 540	cca Pro	atg Met	aat Asn	ctt Leu gga	ttc Phe 545	
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					cct Pro										ctt Leu	2590
					tat Tyr 775										ttc Phe 785	2638
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	cgt tat Arg Tyr		Leu Le				Gly					3358
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ggt Gly	ttt Phe	gac Asp	ttt Phe	atg Met 111	Lys	gtt Val	ggt Gly	gag Glu	gat Asp 111!	Ala	tta Leu	gtt Val	aat Asn	tta Leu 112		3646
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	Ile		att Ile				gctti	tat 1	ttgg	caati	ta aa	aaag	agca	t		4318
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His	Gln	115 Trp	Arg	Met	Leu	Val	120 Ile	Phe	Leu	Val	Ser	125 Ser	Met	Tle	Len	
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Phe Leu Leu Tyr Gly Leu Tyr Ile Ser Gln Asn Gln Glu Ile Val Ala
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Val Arg Gly Leu Leu Leu Asn Ser Gly Asn Leu Thr Ile His Gly Gln
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Tyr Ser Leu Val Gly Phe Gly His His Ile Ile Lys Gln Asp Ser His
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			Arg	85					90					95	
			Glu 100					105					110	_	
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			Met	165					170					175	
			Trp 180 Ile					185					190		_
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			Phe	245					250					255	
			260 Asn					265					270		_
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	290		Phe			295					300			_	
305			Lys		310					315					320
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385			Tyr		390					395					400
			Ile	405					410					415	
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Ala	Gly	Leu 435	Leu	Phe	Pro	Ile		Ala			Thr	Gly 445	Gly	Ser	Ile
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Leu 465	Ile	Leu	Thr	Leu	Val 470			Cys				Ile		Gln	Gly 480
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		515					520					525		Cys	
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Pro 545	Ala	Gly	Asp	Asp		Tyr			Glu	Ala 555	Ile	Glu	Ser	Phe	Ile 560
				565					570					Ile 575	_
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		595					600					605		Lys	_
	610					615					620			Lys	
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	690					695					700			Leu	
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785					790					795				Ile	800
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				165					170			Tyr		175	
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		195					200				-	Gly 205			
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Arg																
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								tta Leu		1777
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								aat Asn		1873
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tta Leu	gat Asp	aaa Lys 880	cct Pro	gat Asp	aca Thr	gca Ala	tat Tyr 885	tca Ser	gac Asp	cct Pro	cac His	tta Leu 890	gat Asp	cat His	ttg Leu	2691
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	Val					Gly				gcc Ala	Leu					3225
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Phe					Phe	Ala		Met	Ser	tta Leu 1						3513
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			Ala					Val		atg Met			Leu	_		3609
		Leu					Lys			cct Pro		Ser	_			3657

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His	gct Ala 1290	ctc Leu	ttt Phe	tca Ser	Phe	ttg Leu 295	cta Leu	ttt Phe	gcc Ala	Asn	ctt Leu L300	ttt Phe	act Thr	atg Met	tat Tyr	3993
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	Glu					Gly					Thr				ctt Leu	4425
Arg					Ile	tct Ser 1455				Glu						4473
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		Lys				tta Leu	Ser							Arg I		4858
			Phe			tta Leu		Ser				_	Tyr			4906
		Phe				ttt Phe	Leu					Ser		_	_	4954
	Leu					tta Leu 1					Thr					5002

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                                             1650
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Asp Ser His Ala Phe Asn Tyr Leu Pro Cys Leu Lys Asn Arg Glu Leu
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Gln Leu Ser Ala Phe Leu Ser Gln Leu Asp Lys Asp Phe Leu Phe Glu
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Thr Ser Glu Gln Ala Trp Ala Ser Leu Ile Leu Ser Met Glu Val Glu
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His Thr Lys Thr Phe Leu Lys Lys Trp Lys Thr Ser Thr His Phe Gln
Lys Asp Val Glu His Ile Val Asp Val Tyr Arg Ile Arg Glu Gln Met
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Gln Ala Glu Gly Ile Arg Lys Ala Arg Gly Leu Met Val Asp Phe Glu
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Ile Val Val Asn Gly Gly Thr Leu Ile Lys Lys Leu Gly Ile Lys Pro
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Gly Pro Gln Met Gly Asp Ile Ile Ser Gln Ile Glu Leu Ala Ile Val
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				165					170	Leu				175	
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			340					345		Ile		_	350		
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Val	Thr	Glu	Leu 420	Leu	Glu	Gln	Phe	Leu 425	Phe	Pro	Arg	Ser	Thr 430	His	Gly

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Ile Glu Lys Ser Gly Phe Phe Val Ile Glu Glu Arg Asp Glu Ile Ile
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Leu Val Lys Met Ile Glu Val Glu Ala Arg Lys Ile Gly Tyr Arg Gln
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Leu Tyr Leu Glu Thr Ala Ser Thr Leu Ser Arg Ala Thr Ala Val Tyr
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<213> Streptococcus

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					Gly 999		_	1056
					tat Tyr 360	_		1104

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PCT/CA99/00114

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Ala Thr Tyr Asp Thr Gly Ser Ser Phe Val Ile Pro His Ile Asp His
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Val Lys Tyr Val Met Gln His Pro Glu Val Arg Pro Asp Val Trp Ser
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100 105 110	Glu Gln
Agn Tou Agn Tla Why Tan Glu 17-1 man his 17-1 - Give Tan Gi	
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60

120

180

240

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Asn Asn Asp Cys Tyr Gln Ala Leu Leu Asn Glu Gln Ser Lys Ala Ile

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(71) Applicant (for all designated States except US): BIOCHEM VACCINS INC. [CA/CA]; 2323 boulevard du Parc Technologique, Sainte-Foy, Québec G1P 4R8 (CA).

(72) Inventors; and

(75) Inventors/Applicants (for US only): BRODEUR, Bernard, R. [CA/CA]; 2401 rue Maritain, Sillery, Québec G1T 1N6 (CA). RIOUX, Clément [CA/CA]; 1012 Jean-Charles Cantin, Ville de Cap Rouge, Québec G1Y 2X1 (CA). BOYER, Martine [CA/CA]; Apt. 204, 25 des Mouettes, Beauport, Québec G1E 7G1 (CA). CHARLEBOIS, Isabelle [CA/CA]; 410 Mirabel, St-Nicolas, Québec G7A 2L5 (CA). HAMEL, Josée [CA/CA]; 2401 rue Maritain, Sillery, Québec G1T 1N6 (CA). MARTIN, Denis [CA/CA]; 4728-G rue Gaboury, St-Augustin-de-Desmaures, Québec G3A 1E9 (CA).

(74) Agents: CÔTE, France et al.; Swabey Ogilvy Renault, Suite 1600, 1981 McGill College Avenue, Montréal, Québec H3A 2Y3 (CA).

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(57) Abstract

Group B streptococcus (GBS) proteins and polynucleotides encoding them are disclosed. Said proteins are antigenic and therefore useful vaccine components for the prophylaxis or therapy of streptococcus infection in animals. Also disclosed are recombinant methods of producing the protein antigens as well as diagnostic assays for detecting streptococcus bacterial infection.

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	International Patent Classification (IPC) or to both national classification	fication and IPC	
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Minimum do IPC 6	cumentation searched (classification system followed by classific CO7K C12N A61K	ation symbols)	
Documentat	tion searched other than minimum documentation to the extent tha	it such documents are included in the fields se	arched
Electronic de	ata base consulted during the international search (name of data	base and, where practical, search terms used)	
C DOCUME	ENTS CONSIDERED TO BE RELEVANT		
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Category	Citation of document, with indication, where appropriate, of the	relevant pessages	
A	MICHEL J L ET AL: "Cloned alph C-protein antigens of group B S elicit protective immunity" INFECTION AND IMMUNITY., vol. 59, no. 6, June 1991 (1991 2023-2028, XP002107260 AMERICAN SOCIETY FOR MICROBIOLO WASHINGTON., US ISSN: 0019-9567 the whole document	treptococci -06), pages	1-48
"A" docume consider "E" earlier of filing of which citation other "P" docume other "P" "P" "P" docume other "P" "P" "P" "P" "P" "P" "P" "P" "P" "P	ent which may throw doubts on priority claim(s) or is cited to establish the publication date of another on or other special reason (as specified) sent referring to an oral disclosure, use, exhibition or means sent published prior to the international filing date but	"T" later document published after the integrated or priority date and not in conflict with cited to understand the principle or the invention "X" document of particular relevance; the cannot be considered novel or cannot involve an inventive step when the document of particular relevance; the cannot be considered to involve an indocument is combined with one or ments, such combination being obvious in the art. "&" document member of the same patent.	ernational filing date the application but eory underlying the claimed invention t be considered to comment is taken alone claimed invention eventive step when the ore other such docu-
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C.(Continu	ation) DOCUMENTS CONSIDERED TO BE RELEVANT	Delevent to chief No.
Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	LACHENAUER C S ET AL: "Cloning and expression in Escherichia coli of a protective surface protein from type V group B Streptococci" ADVANCES IN EXPERIMENTAL MEDICINE AND BIOLOGY, vol. 418, 9 December 1997 (1997-12-09), pages 615-618, XP002107261 SPRING ST., NY, US ISSN: 0065-2598 the whole document	1-48
P,X	DATABASE EMBL [Online] Accession number AF062533, 11 February 1999 (1999-02-11) SPELLERBERG B ET AL: "Streptococcus agalactiae Lmb (lmb) gene, complete cds; and unknown gene." XP002125180 98.9% identity between base 1-2514 of SEQ ID NO 13 and base 988-3501 of AF062533 Translation product (AC: Q9ZHG9) has 98.5% identity in 793 AA overlap with SEQ ID NO 15 and 98.5% identity in 715 AA overlap with SEQ ID 16 & SPELLERBERG B ET AL: "Lmb, a protein with similarities to the LraI adhesin family, mediates attachment of Streptococcus agalactiae to human laminin" INFECTION AND IMMUNITY., vol. 67, no. 2, February 1999 (1999-02), pages 871-878, AMERICAN SOCIETY FOR MICROBIOLOGY. WASHINGTON., US ISSN: 0019-9567	1-10, 16-23,26
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· · · · · · · · · · · · · · · · · · ·	ation) DOCUMENTS CONSIDERED TO BE RELEVANT Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Category °	Citation of document, with indication, where appropriate, or the relevant passages	
X	DATABASE EMBL [Online] Accession Number AF026542, 15 October 1997 (1997-10-15) HYNES W L ET AL: "Streptococcus pyogenes FF22 lantibiotic (scn) gene cluster region containing: scnK, scnR, streptococcin A-FF22 precursor (scnA), scnA1, scnM, scnT, scnF, scnE, scnG genes, complete cds, and tnpA gene, partial cds." XP002125182 88.2% identity between base 2607-2953 of SEQ ID NO 13 and base 10435-10777 of AF026542 Translation product (AC: 031057) has 95.8% identity in 71 AA overlap with SEQ ID NO 17	1-10, 16-23, 26
P,X	DATABASE GENESEQ [Online] Accession Number V52136, 23 October 1998 (1998-10-23) BARASH S C ET AL: "Streptococcus pneumoniae genome fragment SEQ ID NO:3" XP002125183 68.5% identity between base 2539-3319 of SEQ ID NO 37 and base 18492-19271 of V52136 Translation has 74.5% identity in 231 AA overlap with SEQ ID NO 40 & WO 98 18931 A (DOUGHERTY BRIAN A ;HUMAN GENOME SCIENCES INC (US); ROSEN CRAIG A) 7 May 1998 (1998-05-07)	1,3-7,10

ational application No. PCT/CA 99/00114

Box I	Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)
This Inte	ernational Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:
1. X	Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely: Although claims 37-46 are directed to a method of treatment of the
	human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2.	Claims Nos.: because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3.	Claims Nos.:
	because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).
Box II	Observations where unity of invention is lacking (Continuation of item 2 of first sheet)
This Inte	ernational Searching Authority found multiple inventions in this international application, as follows:
	see additional sheet
	As a result of the prior review under R. 40.2(e) PCT, no additional fees are to be refunded.
1.	As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2.	As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
з. 🛛	As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
	11-14,16,24,25,27,28,30,31 (completely), 1-10,15,17-23,26,29,32-48 (all partially) i.e. (group of) inventions 1, 3 and 7
4.	No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
Remari	con Protest X The additional search fees were accompanied by the applicant's protest.
	No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: 1-10,15,17-23,26,29,32-48 (all partially)

An isolated polynucleotide encoding a polypeptide having a sequence selected from the group consisting of SEQ ID 2, SEQ ID NO 3, SEQ ID NO 4, SEQ ID NO 5, SEQ ID NO 6 i.e. the open reading frames of clone 1 (SEQ ID NO 1). Also a vector comprising the polynucleotide, a host cell transformed therewith, an isolated polypeptide encoded by the polynucleotide, a vaccine composition comprising said polypeptide and a polynucleotide having a sequence SEQ ID NO 1.

2. Claims: 1-10,15,17-23,26,29,32-48 (all partially)

Same as invention 1, but directed at polypeptides of clone 2 (SEQ ID 7) with sequences SEQ ID NO 8-12.

3. Claims: 1-10,15,17-23,26,29,32-48 (all partially)

Same as invention 1, but directed at polypeptides of clone 3 (SEQ ID 13) with sequences SEQ ID NO 14-21.

4. Claims: 1-10,15,17-23,26,29,32-48 (all partially)

Same as invention 1, but directed at polypeptides of clone 4 (SEQ ID 22) with sequences SEQ ID NO 23-26.

5. Claims: 1-10,15,17-23,26,29,32-48 (all partially)

Same as invention 1, but directed at polypeptides of clone 5 (SEQ ID 27) with sequences SEQ ID NO 28-31.

6. Claims: 1-10,15,17-23,26,29,32-48 (all partially)

Same as invention 1, but directed at polypeptides of clone 6 (SEQ ID 32) with sequences SEQ ID NO 33-36.

7. Claims: 11-14,16,24,25,27,28,30,31 (all completely), 1-10, 15,17-23,26,29,32-48 (all partially)

Same as invention 1, but directed at polypeptides of clone 7 (SEQ ID 37) with sequences SEQ ID NO 38-41.

information on patent family members

Interr Dal Application No PCT/CA 99/00114

Patent document cited in search report		Publication date		atent family nember(s)	Publication date
WO 9818931	Α	07-05-1998	AU AU EP EP WO	5194598 A 6909098 A 0942983 A 0941335 A 9818930 A	22-05-1998 22-05-1998 22-09-1999 15-09-1999 07-05-1998